

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
4 December 2003 (04.12.2003)

PCT

(10) International Publication Number
WO 03/099276 A1

(51) International Patent Classification⁷: **A61K 31/41**,
31/44, 31/435, C07D 213/14, 213/75, 471/04, 514/300,
514/303, 514/352, 514/406, 514/407, 546/117, 546/119,
546/309, 548/364, 548/369, 548/369

(21) International Application Number: PCT/US03/13893

(22) International Filing Date: 5 May 2003 (05.05.2003)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/379,357 10 May 2002 (10.05.2002) US
60/415,367 2 October 2002 (02.10.2002) US

(71) Applicant (*for all designated States except US*): **BRISTOL-MYERS SQUIBB COMPANY** [US/US]; P.O. BOX 4000, ROUTE 206 and PROVINCELINE ROAD, PRINCETON, NJ 08543-4000 (US).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **QIAO, Jennifer, X.** [CN/US]; 3233 Town Court South, Lawrenceville, NJ 08648 (US). **PINTO, Donald, J.** [US/US]; 425 Brentford Road, Kennett Square, PA 19348 (US). **ORWAT, Michael, J.** [US/US]; 118 S. Colts Neck Way, Hockessin, DE 19707

(US). **HAN, Wei** [CN/US]; 660 ROSALIND RUN, Yardley, PA 19067 (US). **FRIEDRICH, Sarah, R.** [US/US]; 1906 Cherrie Circle, Blue Bell, PA 19422 (US).

(74) Agents: **BELFIELD, Jing, S.** et al.; Bristol-Myers Squibb Company, P.O. Box 4000, Route 206 and Province Line Road, Princeton, NJ 08543-4000 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: 1,1-DISUBSTITUTED CYCLOALKYL DERIVATIVES AS FACTOR XA INHIBITORS

(57) Abstract: The present application describes 1,1-disubstituted cycloalkyl compounds and derivatives thereof, or pharmaceutically acceptable salt forms thereof, which are useful as inhibitors of factor Xa.

WO 03/099276 A1

TITLE

1,1-Disubstituted Cycloalkyl Derivatives As Factor Xa
Inhibitors

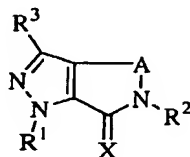
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FIELD OF THE INVENTION

This invention relates generally to 1,1-disubstituted cycloalkyl compounds, which are inhibitors of trypsin-like serine protease enzymes, especially factor Xa,
 10 pharmaceutical compositions containing the same, and methods of using the same as anticoagulant agents for treatment and prevention of thromboembolic disorders.

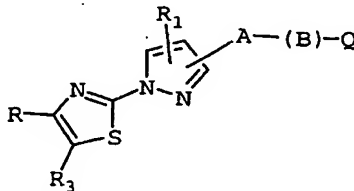
BACKGROUND OF THE INVENTION

15 U.S. Patent Nos. 3,365,459, 3,340,269, and 3,423,414 illustrate anti-inflammatory inhibitors of the following formula:



wherein A is 2-3 carbon atoms, X can be O, and R¹ and R³ can
 20 be substituted or unsubstituted aromatic groups. None of these patents, however, exemplify or suggest compounds of the present invention.

US 5,342,851 depicts thiazole platelet aggregation inhibitors including those of the following formula:



25

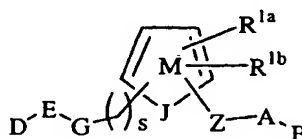
wherein A is a linker, B can be a linker or a ring, Q is a ring or an amino group, R, R₁, and R₃ are a variety of groups. This patent, however, does not exemplify or suggest compounds of the present invention.

WO00/39131 describes heterobicyclic Factor Xa inhibitors of which the following is an example formula:



wherein Z is C or N, G is a mono- or bicyclic group, A is a cyclic moiety and B is a basic group or a cyclic moiety. Compounds specifically described in WO00/39131 are not considered to be part of the present invention.

WO98/28269, WO98/28282, WO99/32454, US 6,020,357, and US 6,271,237 describe Factor Xa inhibitors of the following formula:



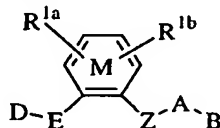
wherein ring M is a heterocycle, Z is a linker, A is a ring, B is a basic or cyclic group, D is a basic moiety, and E is a ring. Compounds specifically described in WO98/28269, WO98/28282, WO99/32454, US 6,020,357, and US 6,271,237 are not considered to be part of the present invention.

WO98/57951 describes Factor Xa inhibitors of the following formula:



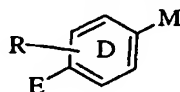
wherein ring M can be a variety of heterocycles and rings D-E represent a heterobicyclic group. Compounds specifically described in WO98/57951 are not considered to be part of the present invention.

WO98/57934 and US 6,060,491 describe Factor Xa inhibitors of the following formula:



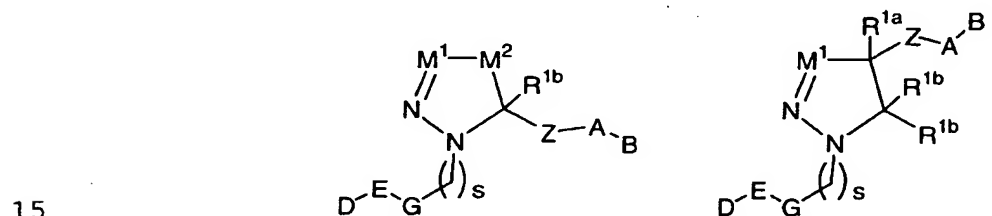
wherein ring M is a 6-membered heteroaryl, Z is a linker, A is a ring, B is a basic or cyclic group, D is a basic moiety, and E is a ring. Compounds specifically described in WO98/57934 and US 6,060,491 are not considered to be
 5 part of the present invention.

WO98/57937 and US 5,998,424 describe Factor Xa inhibitors of the following formula:



wherein ring M is a variety of rings, ring D is an aromatic
 10 ring, and R and E are non-basic groups. Compounds specifically described in WO98/57937 and US 5,998,424 are not considered to be part of the present invention.

WO99/50255 and US 6,191,159 describe pyrazoline and triazoline Factor Xa inhibitors of the following formulas:



Compounds specifically described in WO99/50255 and US 6,191,159 are not considered to be part of the present invention.

WO00/59902 describes Factor Xa inhibitors of the
 20 following formula:

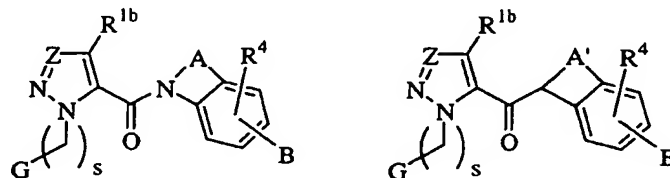


wherein ring M can be a variety of rings all of which are substituted with Z-A-B, Z is a linker, A is a ring, B is a sulfonyl-containing heterobicycle, and rings D-E represent
 25 a heterobicyclic group or a 6-membered ring. Compounds specifically described in WO00/59902 are not considered to be part of the present invention.

WO01/32628 describes cyano-pyrroles, cyano-imidazoles, cyano-pyrazoles, and cyano-triazoles that are Factor Xa

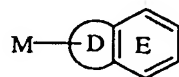
inhibitors. Compounds specifically described in WO01/32628 are not considered to be part of the present invention.

WO01/05784 describes Factor Xa inhibitors of the following formulas:



wherein Z is C or N, G is a mono- or bicyclic ring M, A is a linker, B is a basic or cyclic group. Compounds specifically described in WO01/05784 are not considered to be part of the present invention.

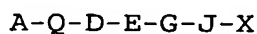
WO00/39108 describes Factor Xa inhibitors of the following formula:



wherein ring M can be a variety of heterocycles and rings D-E represent a heterobicyclic group. Compounds

specifically described in WO00/39108 are not considered to be part of the present invention.

WO01/19798 describes factor Xa inhibitors of the following formula:



wherein A, D, G, and X can be phenyl or heterocycle. However, none of the presently claimed compounds are exemplified or suggested in WO01/19798.

Activated factor Xa, whose major practical role is the generation of thrombin by the limited proteolysis of prothrombin, holds a central position that links the intrinsic and extrinsic activation mechanisms in the final common pathway of blood coagulation. The generation of thrombin, the final serine protease in the pathway to generate a fibrin clot, from its precursor is amplified by formation of prothrombinase complex (factor Xa, factor V, Ca^{2+} and phospholipid). Since it is calculated that one molecule of factor Xa can generate 138 molecules of

thrombin (Elodi, S., Varadi, K.: Optimization of conditions for the catalytic effect of the factor IXa-factor VIII Complex: Probable role of the complex in the amplification of blood coagulation. *Thromb. Res.* **1979**, 15, 617-629),
5 inhibition of factor Xa may be more efficient than inactivation of thrombin in interrupting the blood coagulation system.

Therefore, efficacious and specific inhibitors of factor Xa are needed as potentially valuable therapeutic
10 agents for the treatment of thromboembolic disorders. It is thus desirable to discover new factor Xa inhibitors. In addition, it is also desirable to find new compounds with improved pharmacological characteristics compared with known factor Xa inhibitors. For example, it is preferred
15 to find new compounds with improved factor Xa inhibitory activity and selectivity for factor Xa versus other serine proteases (i.e., trypsin). It is also desirable and preferable to find compounds with advantageous and improved characteristics in one or more of the following categories,
20 but are not limited to: (a) pharmaceutical properties; (b) dosage requirements; (c) factors which decrease blood concentration peak-to-trough characteristics; (d) factors that increase the concentration of active drug at the receptor; (e) factors that decrease the liability for
25 clinical drug-drug interactions; (f) factors that decrease the potential for adverse side-effects; and, (g) factors that improve manufacturing costs or feasibility.

SUMMARY OF THE INVENTION

30 Accordingly, the present invention provides novel 1,1-disubstituted cycloalkyl compounds that are useful as factor Xa inhibitors or pharmaceutically acceptable salts or prodrugs thereof.

The present invention provides pharmaceutical
35 compositions comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of at least

one of the compounds of the present invention or a pharmaceutically acceptable salt or prodrug form thereof.

5 The present invention provides a method for treating thromboembolic disorders comprising administering to a host in need of such treatment a therapeutically effective amount of at least one of the compounds of the present invention or a pharmaceutically acceptable salt or prodrug form thereof.

10 The present invention provides a novel method of treating a patient in need of thromboembolic disorder treatment, comprising: administering a compound of the present invention or a pharmaceutically acceptable salt thereof in an amount effective to treat a thromboembolic disorder.

15 The present invention provides a novel method, comprising: administering a compound of the present invention or a pharmaceutically acceptable salt thereof in an amount effective to treat a thromboembolic disorder.

20 The present invention provides novel compounds for use in therapy.

The present invention provides the use of novel compounds for the manufacture of a medicament for the treatment of a thromboembolic disorder.

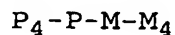
25 These and other objects, which will become apparent during the following detailed description, have been achieved by the inventors' discovery that the presently claimed 1,1-disubstituted cycloalkyl compounds, or pharmaceutically acceptable salt or prodrug forms thereof, are effective factor Xa inhibitors.

30

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

[1] In an embodiment, the present invention provides a novel compound of formula I:

35



I

6

or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

M is a 3-10 membered carbocycle or a 4-10 membered heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, N, and NZ²;

ring M is substituted with 0-3 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

10

P is fused onto ring M and is a 5, 6, or 7 membered carbocycle or a 5, 6, or 7 membered heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

15

ring P is substituted with 0-3 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

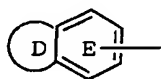
alternatively, ring P is absent and P₄ is directly attached to ring M, provided that when ring P is absent, P₄ and M₄ are attached to the 1,2, 1,3, or 1,4 positions of ring M;

20

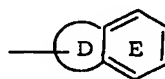
one of P₄ and M₄ is -Z-A-B and the other -G₁-G, provided that P₄ and M₄ are attached to different rings when ring P is present;

25

G is a group of formula IIa or IIb:



IIa



IIb

30

ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;

- 5 E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

alternatively, ring D is absent and ring E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, pyrrolyl, pyrazolyl, imidazolyl, isoxazolyl, oxazolyl, triazolyl, thienyl, and thiazolyl, and ring E is substituted with 1-3 R;

alternatively, ring D is absent and ring E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, pyrrolyl, pyrazolyl, imidazolyl, isoxazolyl, oxazolyl, triazolyl, thienyl, and thiazolyl, and ring E is substituted with 1 R and with a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, wherein the 5-6 membered heterocycle is substituted with 0-2 carbonyls and 1-3 R and there are 0-3 ring double bonds;

- 25 R is selected from H, C₁₋₄ alkyl, F, Cl, Br, I, OH, OCH₃, OCH₂CH₃, OCH(CH₃)₂, OCH₂CH₂CH₃, CN, C(=NR⁸)NR⁷R⁹, NHC(=NR⁸)NR⁷R⁹, ONHC(=NR⁸)NR⁷R⁹, NR⁸CH(=NR⁷), NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂, CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, CH₂CH₂NH₂, CH₂CH₂NH(C₁₋₃ alkyl), CH₂CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tC(O)H, (CR⁸R⁹)_tC(O)R^{2c}, (CR⁸R⁹)_tNR⁷R⁸, (CR⁸R⁹)_tC(O)NR⁷R⁸, (CR⁸R⁹)_tNR⁷C(O)R⁷, (CR⁸R⁹)_tOR³, (CR⁸R⁹)_tS(O)_pNR⁷R⁸, (CR⁸R⁹)_tNR⁷S(O)_pR⁷, (CR⁸R⁹)_tSR³,

$(\text{CR}^8\text{R}^9)_t\text{S}(\text{O})\text{R}^3$, $(\text{CR}^8\text{R}^9)_t\text{S}(\text{O})_2\text{R}^3$, and OCF_3 , provided that $\text{S}(\text{O})_p\text{R}^7$ forms other than $\text{S}(\text{O})_2\text{H}$ or $\text{S}(\text{O})\text{H}$;

alternatively, when 2 R groups are attached to adjacent
 5 atoms, they combine to form methylenedioxy or ethylenedioxy;

A is selected from:

C_{3-10} carbocycle substituted with 0-2 R^4 , and
 10 5-12 membered heterocycle substituted with 0-2 R^4 and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $\text{S}(\text{O})_p$;

B is Y-R^{4a} or X-Y-R^{4a} , provided that Z and B are attached to
 15 different atoms on A and A and R^{4a} or X and R^{4a} are attached to the same atom on Y;

X is selected from $-(\text{CR}^2\text{R}^{2a})_{1-4}-$, $-\text{CR}^2(\text{CR}^2\text{R}^{2b})(\text{CH}_2)_t-$,
 $-\text{C}(\text{O})-$, $-\text{C}(=\text{NR}^{1b})-$, $-\text{CR}^2(\text{NR}^{1b}\text{R}^2)-$, $-\text{CR}^2(\text{OR}^2)-$,
 20 $-\text{CR}^2(\text{SR}^2)-$, $-\text{C}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{C}(\text{O})-$, $-\text{S}(\text{O})-$, $-\text{S}(\text{O})_2-$,
 $-\text{SCR}^2\text{R}^{2a}-$, $-\text{S}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{S}(\text{O})_2\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{S}-$,
 $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})-$, $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})_2-$, $-\text{S}(\text{O})_2\text{NR}^2-$,
 $-\text{S}(\text{O})_2\text{NR}^2\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})_2\text{NR}^2-$, $-\text{NR}^2\text{S}(\text{O})_2-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2\text{S}(\text{O})_2-$, $-\text{NR}^2\text{S}(\text{O})_2\text{CR}^2\text{R}^{2a}-$, $-\text{NR}^2\text{C}(\text{O})-$,
 25 $-\text{C}(\text{O})\text{NR}^2-$, $-\text{NR}^2\text{C}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{C}(\text{O})\text{NR}^2\text{CR}^2\text{R}^{2a}-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2\text{C}(\text{O})-$, $-\text{CR}^2\text{R}^{2a}\text{C}(\text{O})\text{NR}^2-$, NR^2 , $-\text{NR}^2\text{CR}^2\text{R}^{2a}-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2-$, O, $-\text{OCR}^2\text{R}^{2a}-$, and $-\text{CR}^2\text{R}^{2a}\text{O}-$;

Y is a C_{3-10} carbocycle or 3-10 membered heterocycle,
 30 wherein the carbocycle or heterocycle consists of carbon atoms and 0-4 heteroatoms selected from N, O, and $\text{S}(\text{O})_p$, the carbocycle or heterocycle further comprises 0-4 double bonds and 0-2 carbonyl groups, and the carbocycle or heterocycle is substituted with

0-2 R⁴, provided that Y is other than a 1,3-dioxolanyl group;

alternatively, Y is CY¹Y², and Y¹ and Y² are independently
 5 C₁₋₄ alkyl substituted with 0-2 R⁴;

G₁ is absent or is selected from (CR³R^{3a})₁₋₅,

(CR³R^{3a})₀₋₂CR³=CR³(CR³R^{3a})₀₋₂, (CR³R^{3a})₀₋₂C≡C(CR³R^{3a})₀₋₂,
 (CR³R^{3a})_uC(O)(CR³R^{3a})_w, (CR³R^{3a})_uC(O)O(CR³R^{3a})_w,
 10 (CR³R^{3a})_uOC(O)(CR³R^{3a})_w, (CR³R^{3a})_uO(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_w, (CR³R^{3a})_uOC(O)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)O(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)NR^{3b}(CR³R^{3a})_w,
 15 (CR³R^{3a})_uNR^{3b}C(S)NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uS(CR³R^{3a})_w,
 (CR³R^{3a})_uS(O)(CR³R^{3a})_w, (CR³R^{3a})_uS(O)₂(CR³R^{3a})_w,
 (CR³R^{3a})_uS(O)NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uNR^{3b}S(O)₂(CR³R^{3a})_w,
 (CR³R^{3a})_uS(O)₂NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}S(O)₂NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uNR^{3e}(CR³R^{3a})_w,
 20 (CR³R^{3a})_uC(O)(CR³R^{3a})_uC(O)(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(O)(CR³R^{3a})_w,
 (CR³R^{3a})_uC(O)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
 25 (CR³R^{3a})_uNR^{3bb}C(S)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(S)NR^{3b}(CR³R^{3a})_w,
 (CR³R^{3a})_uS(O)NR^{3b}C(O)(CR³R^{3a})_w,
 (CR³R^{3a})_uC(O)NR^{3b}S(O)₂(CR³R^{3a})_w, and
 (CR³R^{3a})_uS(O)₂NR^{3b}C(O)NR^{3b}(CR³R^{3a})_w, wherein u + w total
 30 0, 1, 2, 3, or 4, provided that G₁ does not form a N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

Z is selected from a bond, $-(CR^3R^{3e})_{1-4}-$,

- (CR^3R^{3e})_qO(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qNR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qC(O)(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qC(O)O(CR^3R^{3e})_{q1},
 5 (CR^3R^{3e})_qOC(O)(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qC(O)NR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}C(O)(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qOC(O)O(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qOC(O)NR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}C(O)O(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}C(O)NR^{3b}(CR^3R^{3e})_{q1},
 10 (CR^3R^{3e})_qC(O)(CR^3R^{3e})_qC(O)(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}(CR^3R^{3e})_qC(O)NR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}C(O)(CR^3R^{3e})_qC(O)(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qC(O)(CR^3R^{3e})_qC(O)NR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}C(O)(CR^3R^{3e})_qC(O)NR^{3b}(CR^3R^{3e})_{q1},
 15 (CR^3R^{3e})_qS(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qS(O)(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qS(O)₂(CR^3R^{3e})_{q1}, (CR^3R^{3e})_qSO₂NR^{3b}(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qNR^{3b}SO₂(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qS(O)NR^{3b}C(O)(CR^3R^{3e})_{q1},
 (CR^3R^{3e})_qC(O)NR^{3b}S(O)₂(CR^3R^{3e})_{q1}, and
 20 (CR^3R^{3e})_qNR^{3b}SO₂NR^{3b}(CR^3R^{3e})_{q1}, wherein q + q1 total 0,
 1, 2, 3, or 4, provided that Z does not form a N-S,
 NCH₂N, NCH₂O, or NCH₂S bond with either group to which
 it is attached;

25 provided that:

(a) when ring P is absent and ring M is a pyridyl
 ring, then Z is other than C(O)NHCH₂; and,

(b) when ring P is absent and ring M is a piperazinyl
 ring, then either Z is other than alkylene or A is other
 30 than phenyl;

Z² is selected from H, S(O)₂NHR^{3b}, C(O)R^{3b}, C(O)NHR^{3b},

C(O)OR^{3f}, S(O)R^{3f}, S(O)₂R^{3f}, C₁₋₆ alkyl substituted with

0-2 R^{1a}, C₂₋₆ alkenyl substituted with 0-2 R^{1a}, C₂₋₆ alkynyl substituted with 0-2 R^{1a}, -(C₀₋₄ alkyl)-C₃₋₁₀ carbocycle substituted with 0-3 R^{1a}, and -(C₀₋₄ alkyl)-5-10 membered heterocycle substituted with 0-3 R^{1a} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{1a}, at each occurrence, is selected from H, -(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-CR³R^{1b}R^{1b}, -(CR³R^{3a})_r-O-(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-NR²-(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-S(O)_p-(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-CO₂-(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-C(O)NR²-(CR³R^{3a})_r-R^{1b}, -(CR³R^{3a})_r-C(O)-(CR³R^{3a})_r-R^{1b}, -C₂₋₆ alkenylene-R^{1b}, -C₂₋₆ alkynylene-R^{1b}, and -(CR³R^{3a})_r-C(=NR^{1b})NR³R^{1b}, provided that R^{1a} forms other than an N-halo, N-S, O-O, or N-CN bond;

alternatively, when two R^{1a} groups are attached to adjacent atoms or to the same carbon atom, together with the atoms to which they are attached, they form a 5-7 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p, this ring being substituted with 0-2 R^{4b} and comprising: 0-3 double bonds;

R^{1b} is selected from H, C₁₋₃ alkyl, F, Cl, Br, I, -CN, -NO₂, -CHO, (CF₂)_rCF₃, (CR³R^{3a})_rOR², NR²R^{2a}, C(O)R^{2b}, CO₂R^{2b}, OC(O)R², CH(CH₂OR²)₂, (CF₂)_rCO₂R^{2a}, S(O)_pR^{2b}, NR²(CH₂)_rOR², C(=NR^{2c})NR²R^{2a}, NR²C(O)R^{2b}, NR²C(O)NR²R^{2a}, NR²C(O)₂R^{2a}, OC(O)NR²R^{2a}, C(O)NR²R^{2a}, C(O)NR²(CH₂)_rOR², SO₂NR²R^{2a}, NR²SO₂R², C(O)NR²SO₂R², C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 5-10 membered heterocycle

substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond and
5 provided that $S(O)_p R^2$ forms other than $S(O)_2H$ or $S(O)H$;

R^2 , at each occurrence, is selected from H, CF_3 , C_{1-6} alkyl, benzyl, $-(CH_2)_r-C_{3-10}$ carbocycle substituted with 0-2 R^{4b} , and $-(CH_2)_r-5-10$ membered heterocycle substituted
10 with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2a} , at each occurrence, is selected from H, CF_3 , C_{1-6}
15 alkyl, benzyl, $-(CH_2)_r-C_{3-10}$ carbocycle substituted with 0-2 R^{4b} , and $-(CH_2)_r-5-10$ membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

20 alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1
25 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2b} , at each occurrence, is selected from CF_3 , C_{1-4} alkoxy substituted with 0-2 R^{4b} , C_{1-6} alkyl substituted with
30 0-3 R^{4b} , $-(CH_2)_r-C_{3-10}$ carbocycle substituted with 0-2 R^{4b} , and $-(CH_2)_r-5-10$ membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2c}, at each occurrence, is selected from CF₃, OH, C₁₋₄ alkoxy, C₁₋₆ alkyl, -(CH₂)_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and -(CH₂)_r-5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₆ alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4c}, and -(CR³R^{3a})_r-5-10 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

alternatively, when two R^{2d}'s are attached to the same nitrogen atom, then R^{2d} and R^{2d}, together with the nitrogen atom to which they are attached, combine to form a 5-10 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₆ alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4c}, and -(CR³R^{3a})_r-5-10 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

R^3 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, and phenyl;

5 R^{3a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, and phenyl;

alternatively, R^3 and R^{3a} , together with the nitrogen atom
10 to which they are attached, combine to form a 5 or 6
membered saturated, partially unsaturated, or
unsaturated ring consisting of: carbon atoms, the
nitrogen atom to which R^3 and R^{3a} are attached, and 0-1
additional heteroatoms selected from the group
15 consisting of N, O, and $S(O)_p$;

R^{3b} , at each occurrence, is selected from H, C_{1-6} alkyl
substituted with 0-2 R^{1a} , C_{2-6} alkenyl substituted with
0-2 R^{1a} , C_{2-6} alkynyl substituted with 0-2 R^{1a} , $-(C_{0-4}$
20 alkyl)-5-10 membered carbocycle substituted with 0-3
 R^{1a} , and $-(C_{0-4}$ alkyl)-5-10 membered heterocycle
substituted with 0-3 R^{1a} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and $S(O)_p$;

25

R^{3c} , at each occurrence, is selected from CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, and phenyl;

30 R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, C_{1-4} alkyl-phenyl, and $C(=O)R^{3c}$;

R^{3e} , at each occurrence, is selected from H, $S(O)_2NHR^3$,
 $C(O)R^3$, $C(O)NHR^3$, $C(O)OR^{3f}$, $S(O)R^{3f}$, $S(O)_2R^{3f}$, C_{1-6}
 alkyl substituted with 0-2 R^{1a} , C_{2-6} alkenyl
 substituted with 0-2 R^{1a} , C_{2-6} alkynyl substituted with
 5 0-2 R^{1a} , $-(C_{0-4} \text{ alkyl})$ -5-10 membered carbocycle
 substituted with 0-3 R^{1a} , and $-(C_{0-4} \text{ alkyl})$ - 5-10
 membered heterocycle substituted with 0-3 R^{1a} and
 consisting of: carbon atoms and 1-4 heteroatoms
 selected from the group consisting of N, O, and $S(O)_p$;

10

R^{3f} , at each occurrence, is selected from: C_{1-6} alkyl
 substituted with 0-2 R^{1a} , C_{2-6} alkenyl substituted with
 substituted with 0-2 R^{1a} , C_{2-6} alkynyl substituted with 0-2 R^{1a} , $-(C_{0-4}$
 alkyl)-5-10 membered carbocycle substituted with 0-3
 15 R^{1a} , and $-(C_{0-4} \text{ alkyl})$ -5-10 membered heterocycle
 substituted with 0-3 R^{1a} and consisting of: carbon
 atoms and 1-4 heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$;

20 R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, $-(CH_2)_r$ -3-6 membered
 carbocycle, and $-(CH_2)_r$ -5-6 membered heterocycle
 consisting of: carbon atoms and 1-4 heteroatoms
 25 selected from the group consisting of N, O, and $S(O)_p$;

alternatively, when R^3 and R^{3g} are attached to the same
 carbon atom, they combine with the attached carbon
 atom to form a cyclopropyl group;

30

R^4 , at each occurrence, is selected from H, =O,
 $(CR^3R^{3a})_rOR^2$, F, Cl, Br, I, C_{1-4} alkyl, $(CR^3R^{3a})_rCN$,
 $(CR^3R^{3a})_rNO_2$, $(CR^3R^{3a})_rNR^2R^{2a}$, $(CR^3R^{3a})_rC(O)R^{2c}$,

$(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C}(\text{O})\text{R}^{2b}$, $(\text{CR}^3\text{R}^{3a})_r\text{C}(\text{O})\text{NR}^2\text{R}^{2a}$,
 $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C}(\text{O})\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{C}(=\text{NR}^2)\text{NR}^2\text{R}^{2a}$,
 $(\text{CR}^3\text{R}^{3a})_r\text{C}(=\text{NS}(\text{O})_2\text{R}^{5a})\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C}(=\text{NR}^2)\text{NR}^2\text{R}^{2a}$,
 $(\text{CR}^3\text{R}^{3a})_r\text{C}(\text{O})\text{NR}^2\text{C}(=\text{NR}^2)\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{SO}_2\text{NR}^2\text{R}^{2a}$,
5 $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{SO}_2\text{-C}_{1-4}$ alkyl,
 $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{SO}_2\text{R}^{5a}$, $(\text{CR}^3\text{R}^{3a})_r\text{S}(\text{O})_p\text{R}^{5a}$, $(\text{CR}^3\text{R}^{3a})_r(\text{CF}_2)_r\text{CF}_3$,
 $\text{N}(\text{CH}_2)_r\text{R}^{1b}$, $\text{O}(\text{CH}_2)_r\text{R}^{1b}$, $\text{S}(\text{CH}_2)_r\text{R}^{1b}$, $(\text{CR}^3\text{R}^{3a})_{r-5-6}$
membered carbocycle substituted with 0-1 R^5 , and a
 $(\text{CR}^3\text{R}^{3a})_{r-5-6}$ membered heterocycle substituted with 0-1
10 R^5 and consisting of: carbon atoms and 1-4
heteroatoms selected from the group consisting of N,
O, and $\text{S}(\text{O})_p$;

R^{4a} is selected from C_{1-6} alkyl substituted with 0-2 R^{4c} ,
15 C_{2-6} alkenyl substituted with 0-2 R^{4c} , C_{2-6} alkynyl
substituted with 0-2 R^{4c} , $-(\text{CR}^3\text{R}^{3g})_r\text{-C}_{5-10}$ membered
carbocycle substituted with 0-3 R^{4c} , $-(\text{CR}^3\text{R}^{3g})_{r-5-10}$
membered heterocycle substituted with 0-3 R^{4c} and
consisting of: carbon atoms and 1-4 heteroatoms
20 selected from the group consisting of N, O, and $\text{S}(\text{O})_p$.
 $(\text{CR}^3\text{R}^{3g})_r\text{CN}$, $(\text{CR}^3\text{R}^{3g})_r\text{C}(=\text{NR}^{2d})\text{NR}^{2d}\text{R}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{NR}^{2d}\text{C}(=\text{NR}^{2d})\text{NR}^{2d}\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{NR}^{2d}\text{C}(\text{R}^{2e})(=\text{NR}^{2d})$,
 $(\text{CR}^3\text{R}^{3g})_r\text{NR}^{2d}\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{N}(\rightarrow\text{O})\text{R}^{2d}\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{OR}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{-NR}^{2d}\text{C}(\text{O})\text{R}^{2e}$, $(\text{CR}^3\text{R}^{3g})_r\text{-C}(\text{O})\text{R}^{2e}$,
25 $(\text{CR}^3\text{R}^{3g})_r\text{-OC}(\text{O})\text{R}^{2e}$, $(\text{CR}^3\text{R}^{3g})_r\text{-C}(\text{O})\text{NR}^{2d}\text{R}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{-C}(\text{O})\text{OR}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{-NR}^{2d}\text{C}(\text{O})\text{NR}^{2d}\text{R}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{-OC}(\text{O})\text{NR}^{2d}\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{-NR}^{2d}\text{C}(\text{O})\text{OR}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{-SO}_2\text{NR}^{2d}\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{-NR}^{2d}\text{SO}_2\text{NR}^{2d}\text{R}^{2d}$,
 $(\text{CR}^3\text{R}^{3g})_r\text{-C}(\text{O})\text{NR}^{2d}\text{SO}_2\text{R}^{2d}$, $(\text{CR}^3\text{R}^{3g})_r\text{-NR}^{2d}\text{SO}_2\text{R}^{2d}$, and
30 $(\text{CR}^3\text{R}^{3g})_r\text{-S}(\text{O})_p\text{R}^{2d}$, provided that $\text{S}(\text{O})_p\text{R}^{2d}$ forms other
than $\text{S}(\text{O})_2\text{H}$ or $\text{S}(\text{O})\text{H}$ and further provided that R^{4a} is
other than a hydroxamic acid;

R^{4b} , at each occurrence, is selected from H, =O, $(CH_2)_rOR^3$,
 $(CH_2)_rF$, $(CH_2)_rCl$, $(CH_2)_rBr$, $(CH_2)_rI$, C_{1-4} alkyl,
 $(CH_2)_rCN$, $(CH_2)_rNO_2$, $(CH_2)_rNR^3R^{3a}$, $(CH_2)_rC(O)R^3$,
5 $(CH_2)_rC(O)OR^{3c}$, $(CH_2)_rNR^3C(O)R^{3a}$, $(CH_2)_rC(O)NR^3R^{3a}$,
 $(CH_2)_rNR^3C(O)NR^3R^{3a}$, $(CH_2)_rC(=NR^3)NR^3R^{3a}$,
 $(CH_2)_rNR^3C(=NR^3)NR^3R^{3a}$, $(CH_2)_rSO_2NR^3R^{3a}$,
 $(CH_2)_rNR^3SO_2NR^3R^{3a}$, $(CH_2)_rNR^3SO_2-C_{1-4}$ alkyl,
 $(CH_2)_rNR^3SO_2CF_3$, $(CH_2)_rNR^3SO_2$ -phenyl, $(CH_2)_rS(O)_pCF_3$,
10 $(CH_2)_rS(O)_p-C_{1-4}$ alkyl, $(CH_2)_rS(O)_p$ -phenyl, and
 $(CH_2)_r(CF_2)_rCF_3$;

R^{4c} , at each occurrence, is selected from =O, $(CR^3R^{3a})_rOR^2$,
 $(CR^3R^{3a})_rF$, $(CR^3R^{3a})_rBr$, $(CR^3R^{3a})_rCl$, $(CR^3R^{3a})_rCF_3$, C_{1-4}
15 alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, $(CR^3R^{3a})_rCN$,
 $(CR^3R^{3a})_rNO_2$, $(CR^3R^{3a})_rNR^2R^{2a}$, $(CR^3R^{3a})_rN(\rightarrow O)R^2R^{2a}$,
 $(CR^3R^{3a})_rC(O)R^{2c}$, $(CR^3R^{3a})_rNR^2C(O)R^{2b}$,
 $(CR^3R^{3a})_rC(O)NR^2R^{2a}$, $(CR^3R^{3a})_rN=CHOR^3$,
 $(CR^3R^{3a})_rC(O)NR^2(CH_2)_2NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2C(O)NR^2R^{2a}$,
20 $(CR^3R^{3a})_rC(=NR^2)NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2C(=NR^2)NR^2R^{2a}$,
 $(CR^3R^{3a})_rSO_2NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2SO_2NR^2R^{2a}$,
 $(CR^3R^{3a})_rC(O)NR^2SO_2-C_{1-4}$ alkyl, $(CR^3R^{3a})_rNR^2SO_2R^{5a}$,
 $(CR^3R^{3a})_rS(O)_pR^{5a}$, $(CF_2)_rCF_3$, $(CR^3R^{3a})_rC_{3-10}$ carbocycle
substituted with 0-2 R^{4b} , and $(CR^3R^{3a})_r4-10$ membered
25 heterocycle substituted with 0-2 R^{4b} and consisting of
carbon atoms and from 1-4 heteroatoms selected from
the group consisting of N, O, and $S(O)_p$;

R^5 , at each occurrence, is selected from H, C_{1-6} alkyl, =O,
30 $(CH_2)_rOR^3$, F, Cl, Br, I, -CN, NO_2 , $(CH_2)_rNR^3R^{3a}$,
 $(CH_2)_rC(O)R^3$, $(CH_2)_rC(O)OR^{3c}$, $(CH_2)_rNR^3C(O)R^{3a}$,
 $(CH_2)_rC(O)NR^3R^{3a}$, $(CH_2)_rNR^3C(O)NR^3R^{3a}$, $(CH_2)_rCH(=NOR^{3d})$,

$(\text{CH}_2)_r\text{C}(=\text{NR}^3)\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{C}(=\text{NR}^3)\text{NR}^3\text{R}^{3a}$,
 $(\text{CH}_2)_r\text{SO}_2\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{-C}_{1-4}$
 alkyl, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{CF}_3$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{-phenyl}$,
 $(\text{CH}_2)_r\text{S(O)}_p\text{CF}_3$, $(\text{CH}_2)_r\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $(\text{CH}_2)_r\text{S(O)}_p\text{-}$
 5 phenyl, $(\text{CF}_2)_r\text{CF}_3$, phenyl substituted with 0-2 R^6 ,
 naphthyl substituted with 0-2 R^6 , and benzyl
 substituted with 0-2 R^6 ;

R^{5a} , at each occurrence, is selected from C_{1-6} alkyl,
 10 $(\text{CH}_2)_r\text{OR}^3$, $(\text{CH}_2)_r\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{C(O)}\text{R}^3$, $(\text{CH}_2)_r\text{C(O)}\text{OR}^{3c}$,
 $(\text{CH}_2)_r\text{NR}^3\text{C(O)}\text{R}^{3a}$, $(\text{CH}_2)_r\text{C(O)}\text{NR}^3\text{R}^{3a}$, $(\text{CF}_2)_r\text{CF}_3$, phenyl
 substituted with 0-2 R^6 , naphthyl substituted with 0-2
 R^6 , and benzyl substituted with 0-2 R^6 , provided that
 R^{5a} does not form a S-N or $\text{S(O)}_p\text{-C(O)}$ bond;

15 R^6 , at each occurrence, is selected from H, OH, $(\text{CH}_2)_r\text{OR}^2$,
 halo, C_{1-4} alkyl, -CN, NO_2 , $(\text{CH}_2)_r\text{NR}^2\text{R}^{2a}$, $(\text{CH}_2)_r\text{C(O)}\text{R}^{2b}$,
 $\text{NR}^2\text{C(O)}\text{R}^{2b}$, $\text{NR}^2\text{C(O)}\text{NR}^2\text{R}^{2a}$, C(=NH)NH_2 , NHC(=NH)NH_2 ,
 $\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{NR}^2\text{SO}_2\text{NR}^2\text{R}^{2a}$, and $\text{NR}^2\text{SO}_2\text{C}_{1-4}$ alkyl;

20 R^7 , at each occurrence, is selected from H, OH, C_{1-6} alkyl,
 C_{1-6} alkyl-C(O)-, C_{1-6} alkyl-O-, $(\text{CH}_2)_n\text{-phenyl}$, C_{1-4}
 alkyl-OC(O)-, C_{6-10} aryl-O-, C_{6-10} aryl-OC(O)-, C_{6-10}
 aryl- $\text{CH}_2\text{-C(O)-}$, C_{1-4} alkyl-C(O)O- C_{1-4} alkyl-OC(O)-,
 25 C_{6-10} aryl-C(O)O- C_{1-4} alkyl-OC(O)-,
 C_{1-6} alkyl- $\text{NH}_2\text{-C(O)-}$, phenyl- $\text{NH}_2\text{-C(O)-}$, and phenyl C_{1-4}
 alkyl-C(O)-;

R^8 , at each occurrence, is selected from H, C_{1-6} alkyl, and
 30 $(\text{CH}_2)_n\text{-phenyl}$;

alternatively, R^7 and R^8 , when attached to the same
 nitrogen, combine to form a 5-10 membered heterocyclic

ring consisting of carbon atoms and 0-2 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

5 R⁹, at each occurrence, is selected from H, C₁₋₆ alkyl, and (CH₂)_n-phenyl;

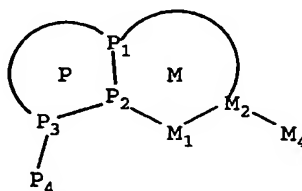
n, at each occurrence, is selected from 0, 1, 2, and 3;

10 p, at each occurrence, is selected from 0, 1, and 2;

r, at each occurrence, is selected from 0, 1, 2, 3, 4, 5, and 6; and,

15 t, at each occurrence, is selected from 0, 1, 2, and 3.

[2] In a preferred embodiment, the present invention provides a novel compound of Formula II:



II

or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

25 ring M, including P₁, P₂, M₁, and M₂, is a 5, 6, or 7 membered carbocycle or a 5, 6, or 7 membered heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, N, and NZ²;

30 ring M is substituted with 0-2 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

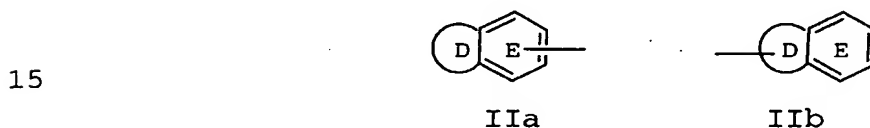
ring P, including P₁, P₂, and P₃, is a 5 or 6 membered aromatic heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

5 alternatively, ring P, including P₁, P₂, and P₃, is a 5 or 6 membered dihydro-aromatic heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

10 ring P is substituted with 0-2 R^{1a};

one of P₄ and M₄ is -Z-A-B and the other -G₁-G;

G is a group of formula IIa or IIb:



ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

20

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;

25 E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

alternatively, ring D is absent, and ring E is selected from phenyl, pyridyl, pyrimidyl, and thienyl, and ring E is substituted with 1-3 R;

30

alternatively, ring D is absent, ring E is selected from phenyl, pyridyl, and thienyl, and ring E is substituted with 1 R and substituted with a 5-6

membered heterocycle consisting of: carbon atoms and
 1-4 heteroatoms selected from the group consisting of
 N, O, and S(O)_p, wherein the 5-6 membered heterocycle
 is substituted with 0-2 carbonyls and 1-3 R and there
 5 are 0-3 ring double bonds;

R is selected from H, C₁₋₄ alkyl, F, Cl, OH, OCH₃, OCH₂CH₃,
 OCH(CH₃)₂, CN, C(=NH)NH₂, C(=NH)NHOH, C(=NH)NHOCH₃,
 NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂,
 10 CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tNR⁷R⁸,
 C(O)NR⁷R⁸, CH₂C(O)NR⁷R⁸, S(O)_pNR⁷R⁸, CH₂S(O)_pNR⁷R⁸,
 SO₂R³, and OCF₃;

alternatively, when 2 R groups are attached to adjacent
 15 atoms, they combine to form methylenedioxy or
 ethylenedioxy;

A is selected from:

C₅₋₁₀ carbocycle substituted with 0-2 R⁴, and
 20 5-10 membered heterocycle substituted with 0-2 R⁴ and
 consisting of: carbon atoms and 1-4 heteroatoms selected
 from the group consisting of N, O, and S(O)_p;

X is selected from -(CR²R^{2a})₁₋₄-, -C(O)-, -C(O)CR²R^{2a}-,
 25 -CR²R^{2a}C(O)-, -S(O)₂-, -S(O)₂CR²R^{2a}-, -CR²R^{2a}S(O)₂-,
 -NR²S(O)₂-, -S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR²,
 -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

Y is a C₃₋₇ monocyclic carbocycle or 3-7 membered monocyclic
 30 heterocycle, wherein the carbocycle or heterocycle
 consists of: carbon atoms and 0-2 heteroatoms
 selected from N, O, and S(O)_p, the carbocycle or
 heterocycle further comprises 0-2 double bonds and 0-2

carbonyl groups, and the carbocycle or heterocycle is substituted with 0-2 R⁴;

alternatively, Y is CY¹Y², and Y¹ and Y² are independently
5 C₁₋₃ alkyl substituted with 0-1 R⁴;

Z is selected from a bond, CH₂, CH₂CH₂, CH₂O, OCH₂, C(O),
NH, CH₂NH, NHCH₂, CH₂C(O), C(O)CH₂, C(O)NH, NHC(O),
NHC(O)CH₂C(O)NH, S(O)₂, CH₂S(O)₂, S(O)₂(CH₂), SO₂NH,
10 and NHSO₂, provided that Z does not form a N-S, NCH₂N,
NCH₂O, or NCH₂S bond with either group to which it is
attached;

Z² is selected from H, C₁₋₄ alkyl, phenyl, benzyl, C(O)R^{3b},
15 S(O)R^{3f}, and S(O)₂R^{3f};

R^{1a}, at each occurrence, is selected from H, -(CH₂)_r-R^{1b},
-(CH(CH₃))_r-R^{1b}, -(C(CH₃)₂)_r-R^{1b}, -O-(CR^{3R3a})_r-R^{1b},
-NR²-(CR^{3R3a})_r-R^{1b}, and -S-(CR^{3R3a})_r-R^{1b}, provided that
20 R^{1a} forms other than an N-halo, N-S, O-O, or N-CN bond;

alternatively, when two R^{1a} groups are attached to adjacent
atoms or to the same carbon atom, together with the
atoms to which they are attached, they form a 5-7
25 membered ring consisting of: carbon atoms and 0-2
heteroatoms selected from the group consisting of N,
O, and S(O)_p, this ring being substituted with 0-2 R^{4b}
and comprising: 0-3 double ring bonds;

30 R^{1b} is selected from H, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, F,
Cl, Br, I, -CN, -CHO, CF₃, OR², NR²R^{2a}, C(O)R^{2b}, CO₂R^{2b},
OC(O)R², CO₂R^{2a}, S(O)_pR^{2b}, NR²(CH₂)_rOR², NR²C(O)R^{2b},
NR²C(O)NHR², NR²C(O)₂R^{2a}, OC(O)NR²R^{2a}, C(O)NR²R^{2a},

$C(O)NR^2(CH_2)_rOR^2$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{5-6} carbocycle
 substituted with 0-2 R^{4b} , and 5-6 membered heterocycle
 substituted with 0-2 R^{4b} and consisting of carbon atoms
 and from 1-4 heteroatoms selected from the group
 5 consisting of N, O, and $S(O)_p$, provided that R^{1b} forms
 other than an O-O, N-halo, N-S, or N-CN bond and
 provided that $S(O)_pR^2$ forms other than $S(O)_2H$ or $S(O)H$;

R^2 , at each occurrence, is selected from H, CF_3 , CH_3 ,
 10 CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle
 substituted with 0-2 R^{4b} , a C_{5-6} carbocycle- CH_2 -
 substituted with 0-2 R^{4b} , and 5-6 membered heterocycle
 substituted with 0-2 R^{4b} and consisting of: carbon
 15 atoms and 1-4 heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$;

R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 ,
 CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 20 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle
 substituted with 0-2 R^{4b} , and 5-6 membered heterocycle
 substituted with 0-2 R^{4b} and consisting of: carbon
 atoms and 1-4 heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$;

25 alternatively, R^2 and R^{2a} , together with the nitrogen atom
 to which they are attached, combine to form a 5 or 6
 membered saturated, partially saturated or unsaturated
 ring substituted with 0-2 R^{4b} and consisting of: 0-1
 30 additional heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$;

R^{2b} , at each occurrence, is selected from CF_3 , C_{1-4} alkoxy,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$,

CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₅-6 carbocycle substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2c}, at each occurrence, is selected from CF₃, OH, C₁-4 alkoxy, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₅-6 carbocycle substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁-4 alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃-6 carbocycle substituted with 0-2 R^{4c}, and -(CR³R^{3a})_r-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

alternatively, when two R^{2d}'s are attached to the same nitrogen atom, then R^{2d} and R^{2d}, together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁-4 alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃-6 carbocycle

substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_x$ -5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2e}
5 forms other than a $C(O)$ -halo or $C(O)-S(O)_p$ moiety;

R^3 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

10 R^{3a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

alternatively, R^3 and R^{3a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6
15 membered saturated, partially unsaturated, or unsaturated ring consisting of: carbon atoms and the nitrogen atom to which R^3 and R^{3a} are attached;

R^{3c} , at each occurrence, is selected from CH_3 , CH_2CH_3 ,
20 $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, CH_2 -phenyl, CH_2CH_2 -phenyl, and $C(=O)R^{3c}$;
25

R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, cyclopropyl-methyl, benzyl, and phenyl;

30 alternatively, when R^3 and R^{3g} are attached to the same carbon atom, they combine with the attached carbon atom to form a cyclopropyl group;

- R^4 , at each occurrence, is selected from H, =O, OR^2 , CH_2OR^2 , $(CH_2)_2OR^2$, F, Cl, Br, I, C_{1-4} alkyl, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $(CH_2)_2NR^2R^{2a}$, $C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , 5-6 membered carbocycle substituted with 0-1 R^5 , and a 5-6 membered heterocycle substituted with 0-1 R^5 and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 10 R^{4b} , at each occurrence, is selected from H, =O, OR^3 , CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, -CN, NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2-C(O)R^3$, $C(O)OR^{3c}$, $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $CH_2NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$,
 15 $CH_2C(O)NR^3R^{3a}$, $NR^3C(O)NR^3R^{3a}$, $CH_2NR^3C(O)NR^3R^{3a}$, $C(=NR^3)NR^3R^{3a}$, $CH_2C(=NR^3)NR^3R^{3a}$, $NR^3C(=NR^3)NR^3R^{3a}$, $CH_2NR^3C(=NR^3)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $CH_2SO_2NR^3R^{3a}$, $NR^3SO_2NR^3R^{3a}$, $CH_2NR^3SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, $CH_2NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, $CH_2NR^3SO_2CF_3$,
 20 NR^3SO_2 -phenyl, $CH_2NR^3SO_2$ -phenyl, $S(O)_pCF_3$, $CH_2S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $CH_2S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, $CH_2S(O)_p$ -phenyl, CF_3 , and CH_2-CF_3 ;
- R^{4c} , at each occurrence, is selected from =O, $(CR^3R^{3a})_rOR^2$,
 25 $(CR^3R^{3a})_rF$, $(CR^3R^{3a})_rBr$, $(CR^3R^{3a})_rCl$, $(CR^3R^{3a})_rCF_3$, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, $(CR^3R^{3a})_rCN$, $(CR^3R^{3a})_rNO_2$, $(CR^3R^{3a})_rNR^2R^{2a}$, $(CR^3R^{3a})_rN(\rightarrow O)R^2R^{2a}$, $(CR^3R^{3a})_rC(O)R^{2c}$, $(CR^3R^{3a})_rNR^2C(O)R^{2b}$, $(CR^3R^{3a})_rC(O)NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2C(O)NR^2R^{2a}$,
 30 $(CR^3R^{3a})_rSO_2NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2SO_2NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2SO_2R^{5a}$, $(CR^3R^{3a})_rS(O)_pR^{5a}$, $(CF_2)_rCF_3$, $(CR^3R^{3a})_rC_{3-10}$ carbocycle substituted with 0-2 R^{4b} , and

(CR³R^{3a})_r 5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

5

R⁵, at each occurrence, is selected from H, =O, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, OR³, CH₂OR³, F, Cl, -CN, NO₂, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, CH₂C(O)R³, C(O)OR^{3c}, CH₂C(O)OR^{3c}, NR³C(O)R^{3a}, C(O)NR³R^{3a}, NR³C(O)NR³R^{3a}, CH(=NOR^{3d}), C(=NR³)NR³R^{3a}, NR³C(=NR³)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, NR³SO₂CF₃, NR³SO₂-phenyl, S(O)_pCF₃, S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl, CF₃, phenyl substituted with 0-2 R⁶, naphthyl substituted with 0-2 R⁶, and benzyl substituted with 0-2 R⁶;

15

R⁶, at each occurrence, is selected from H, OH, OR², F, Cl, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, -CN, NO₂, NR²R^{2a}, CH₂NR²R^{2a}, C(O)R^{2b}, CH₂C(O)R^{2b}, NR²C(O)R^{2b}, NR²C(O)NR²R^{2a}, C(=NH)NH₂, NHC(=NH)NH₂, SO₂NR²R^{2a}, NR²SO₂NR²R^{2a}, and NR²SO₂C₁₋₄ alkyl; and,

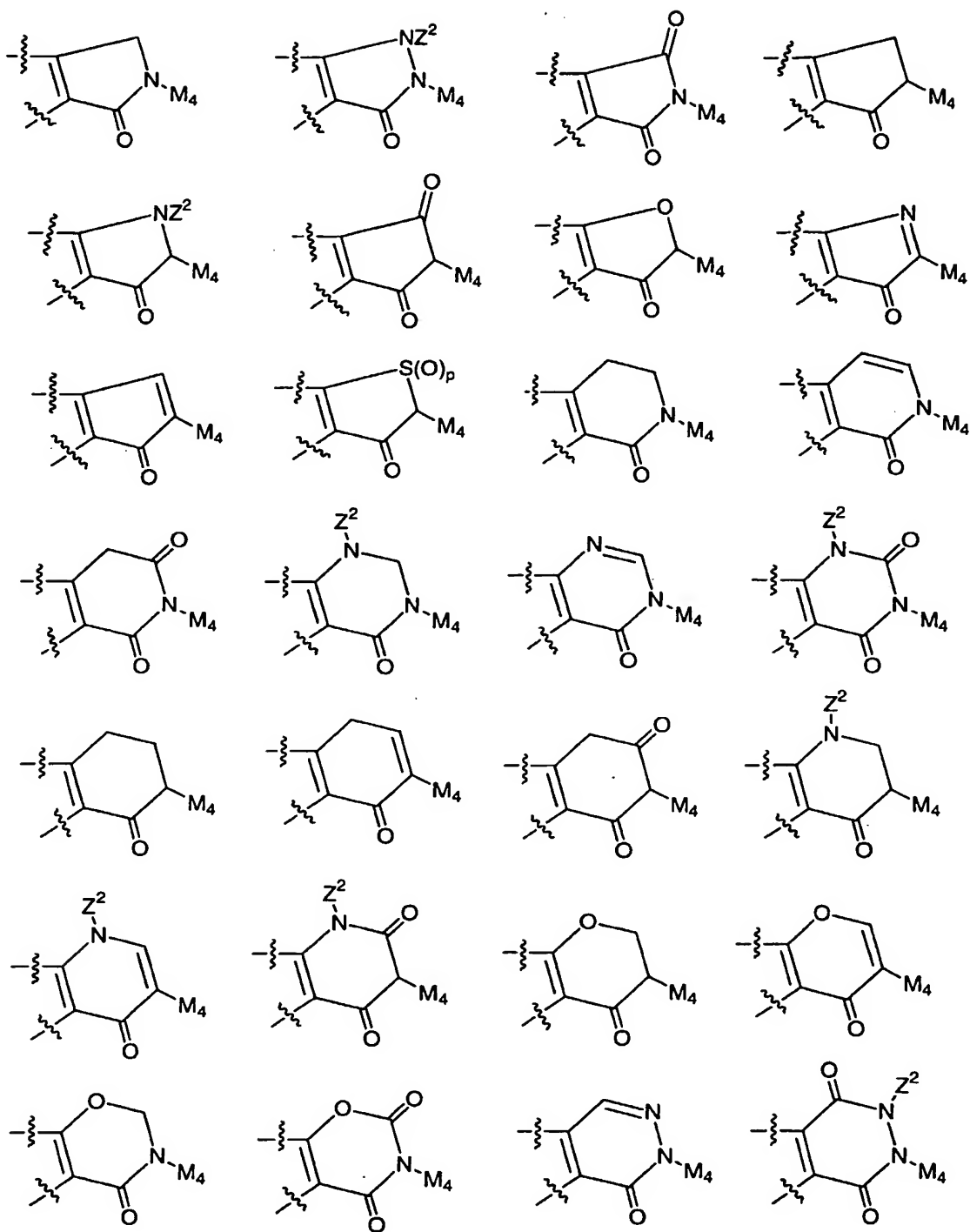
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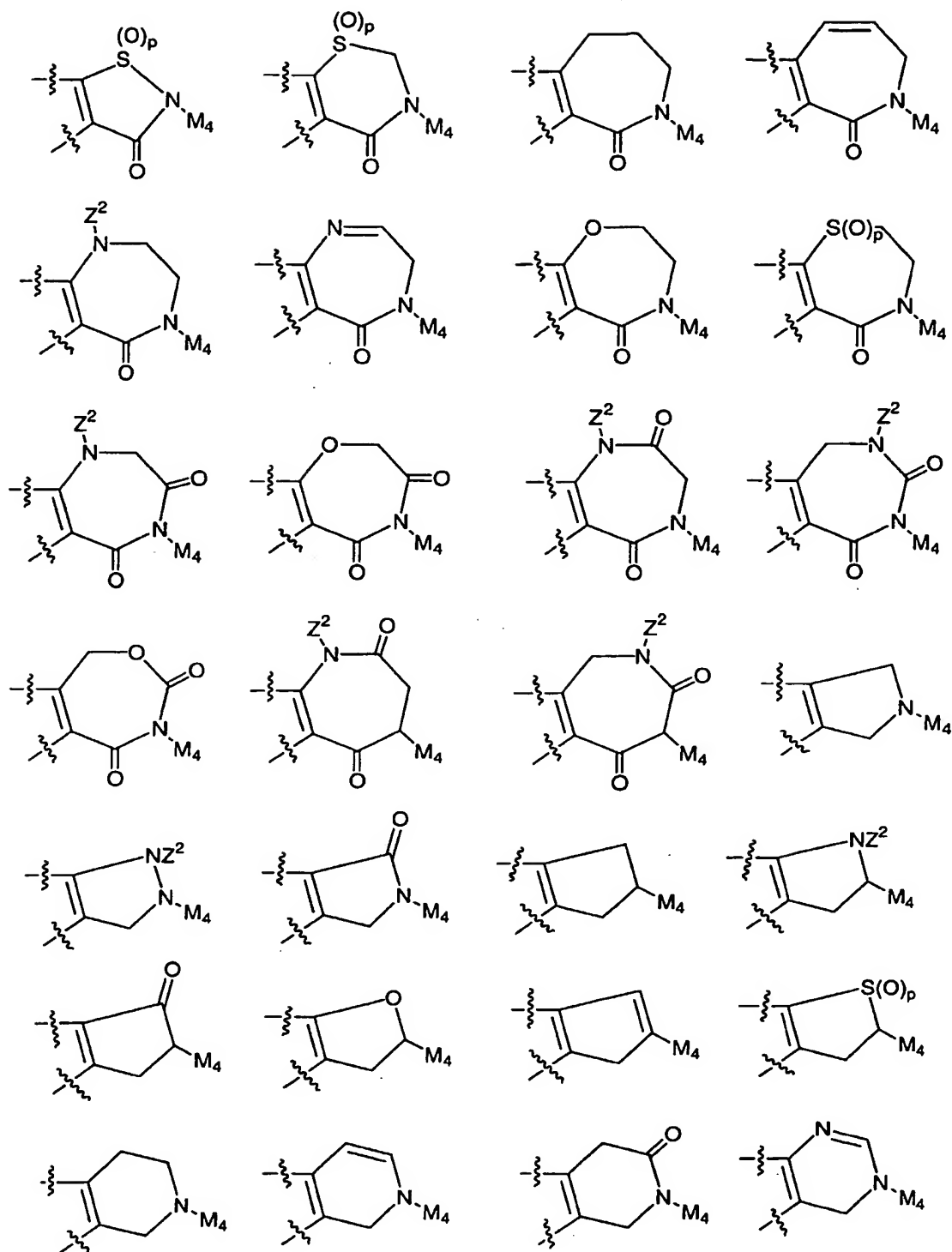
r, at each occurrence, is selected from 0, 1, 2, and 3.

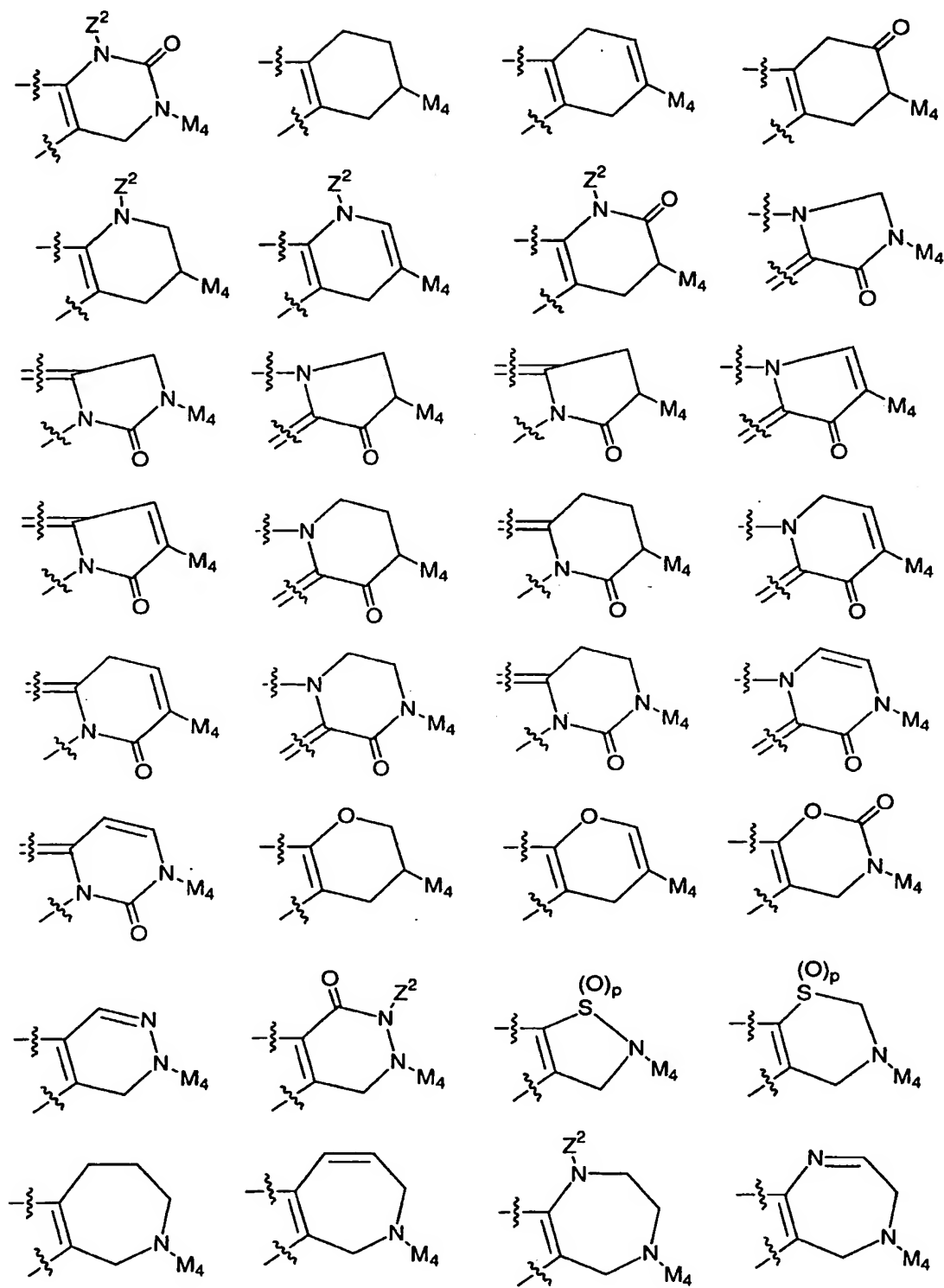
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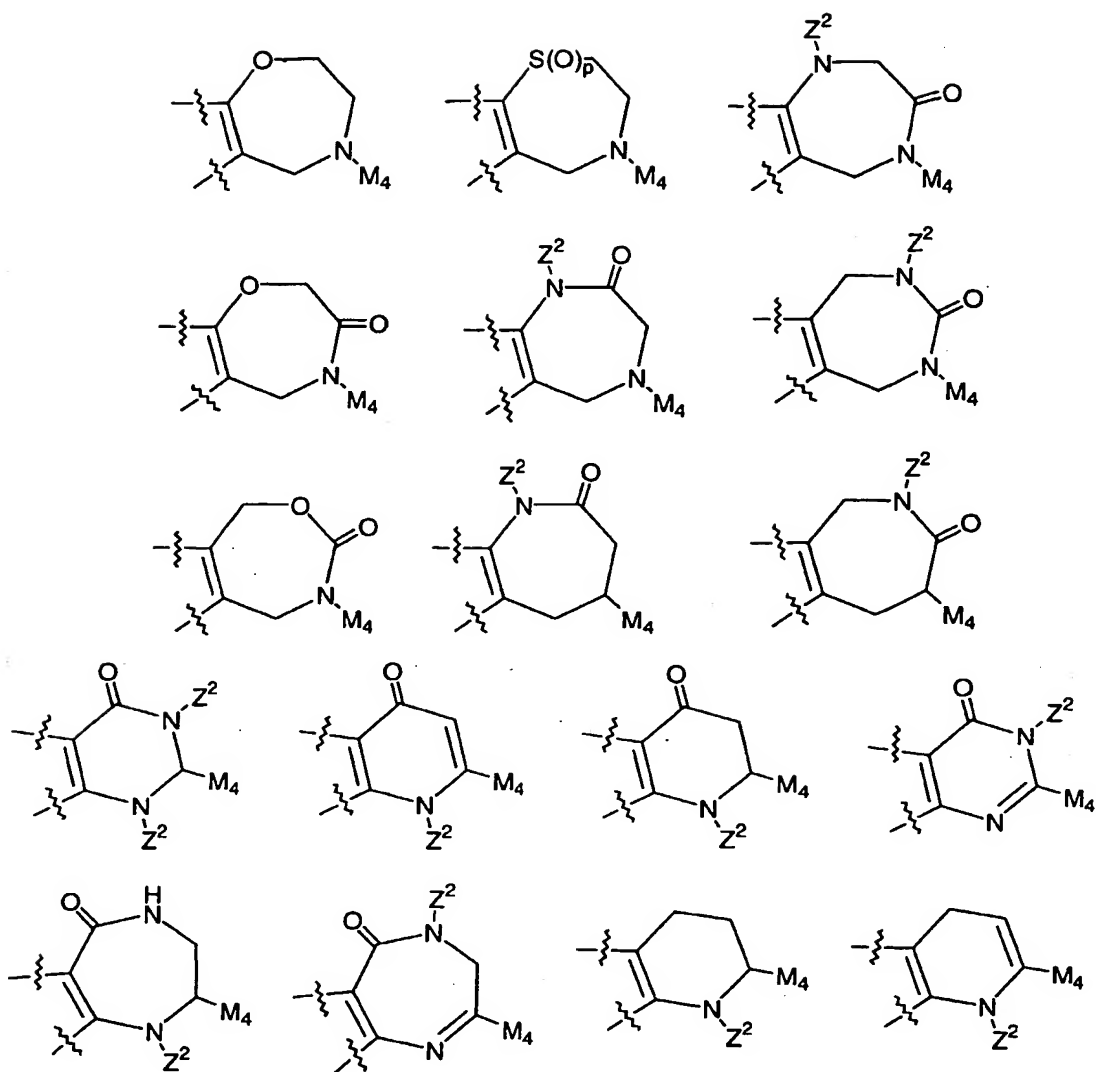
[3] In another preferred embodiment, the present invention provides a novel compound, wherein:

30 ring M is substituted with 0-2 R^{1a} and is selected from the group:



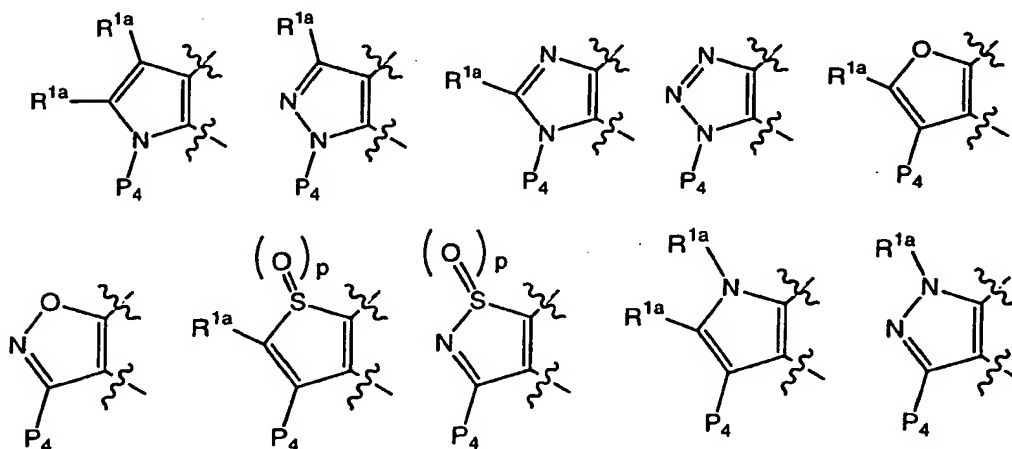


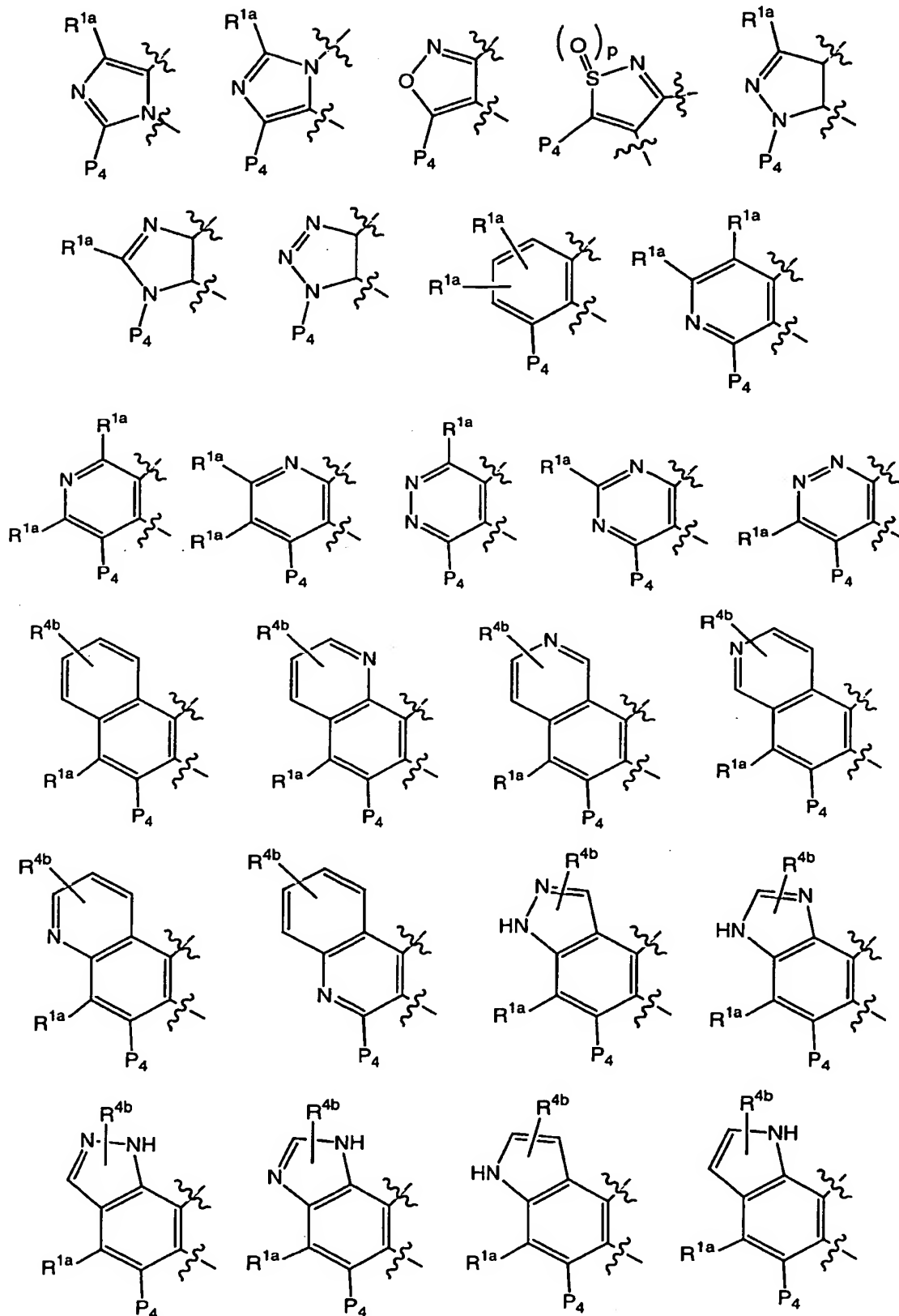


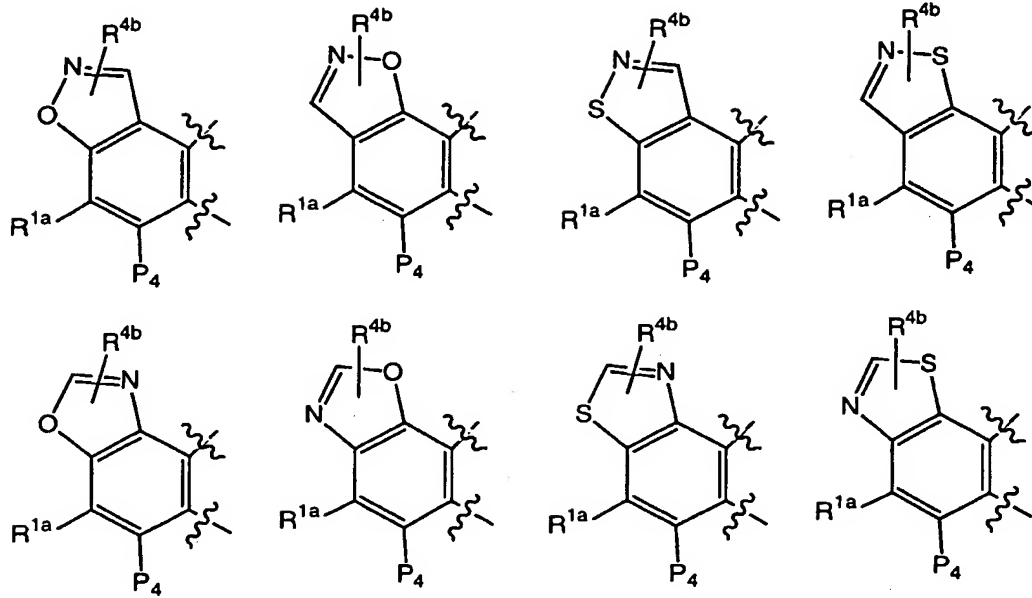


ring P, including P₁, P₂, P₃, and P₄ is selected from group:

5



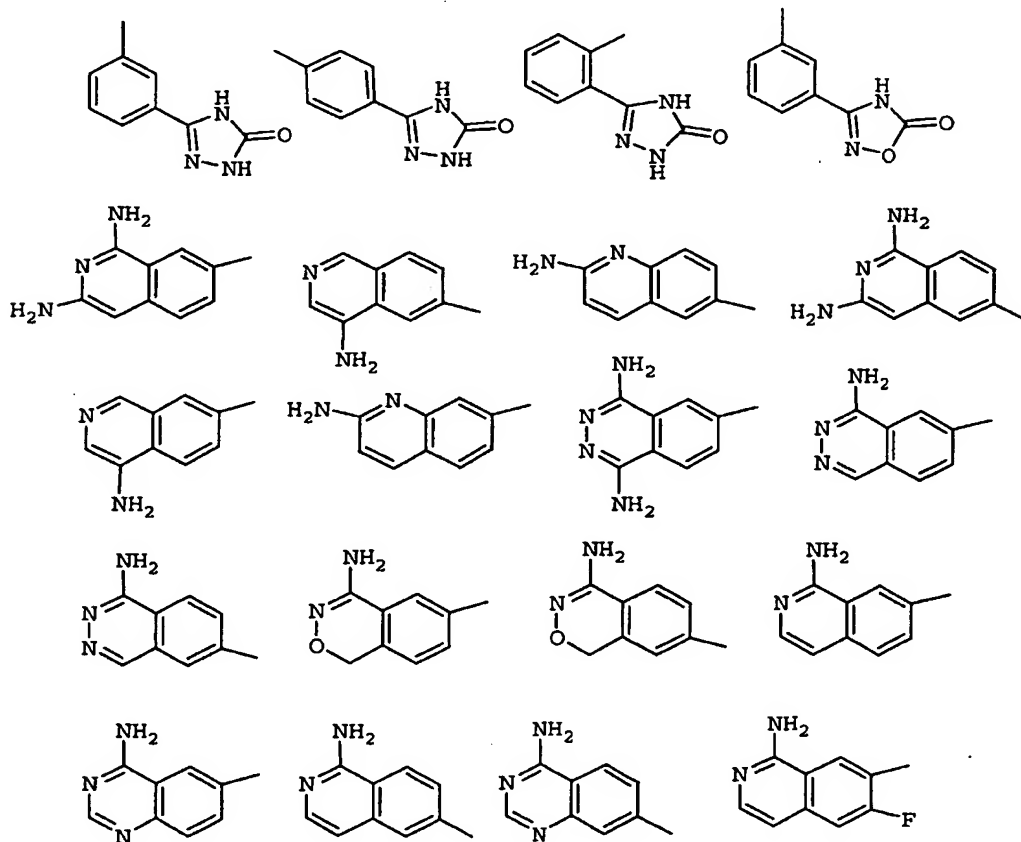


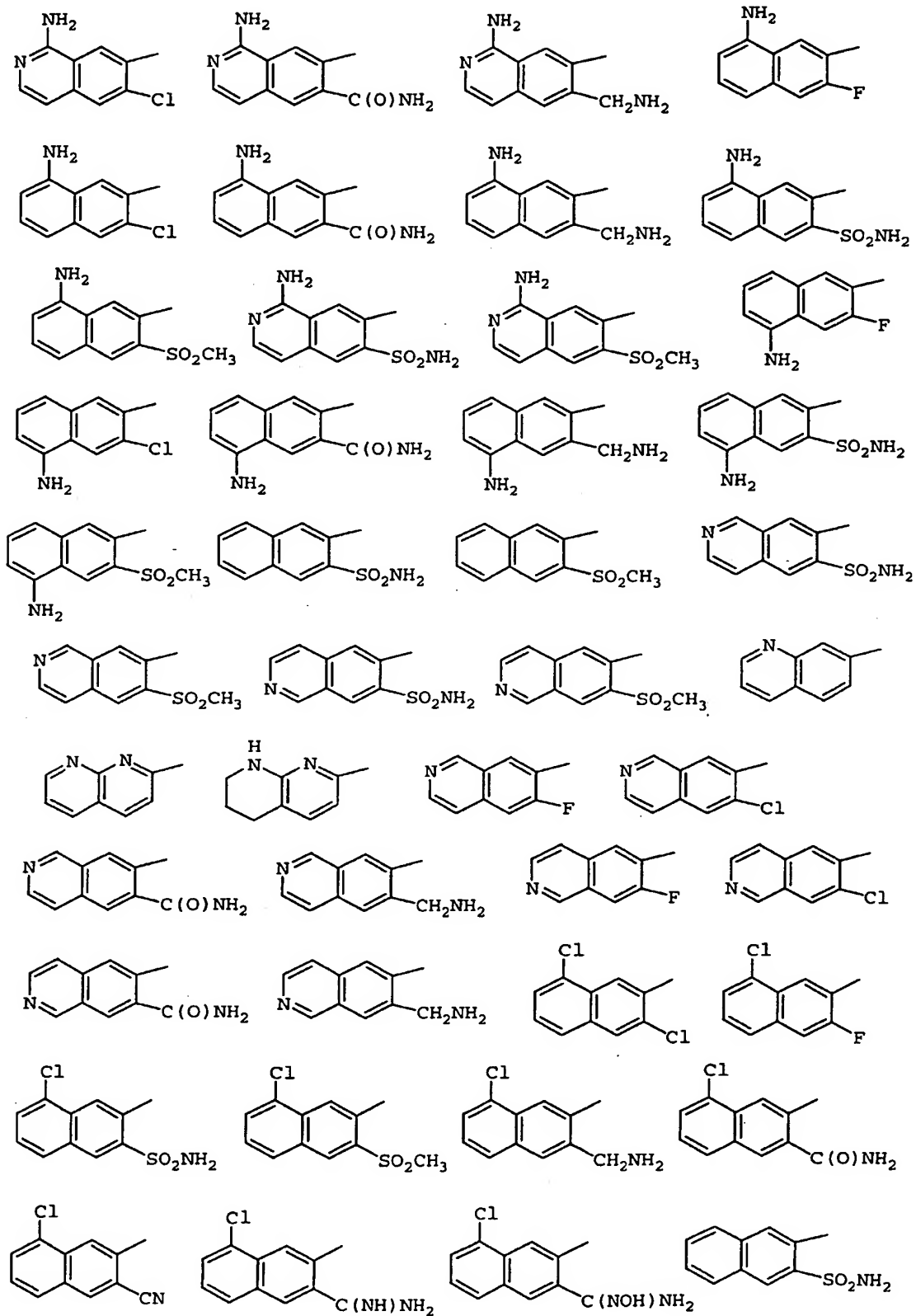


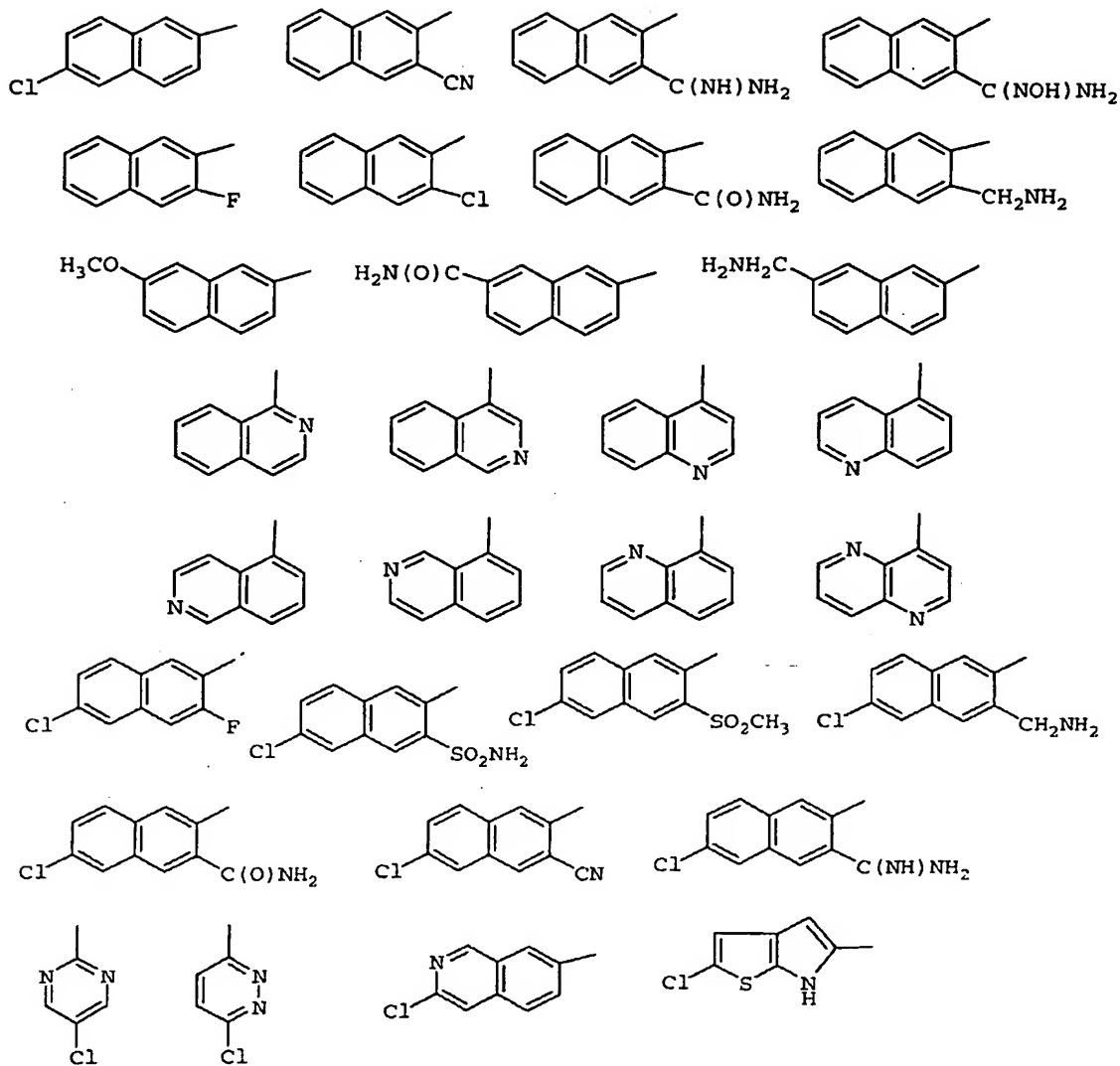
G is selected from the group:

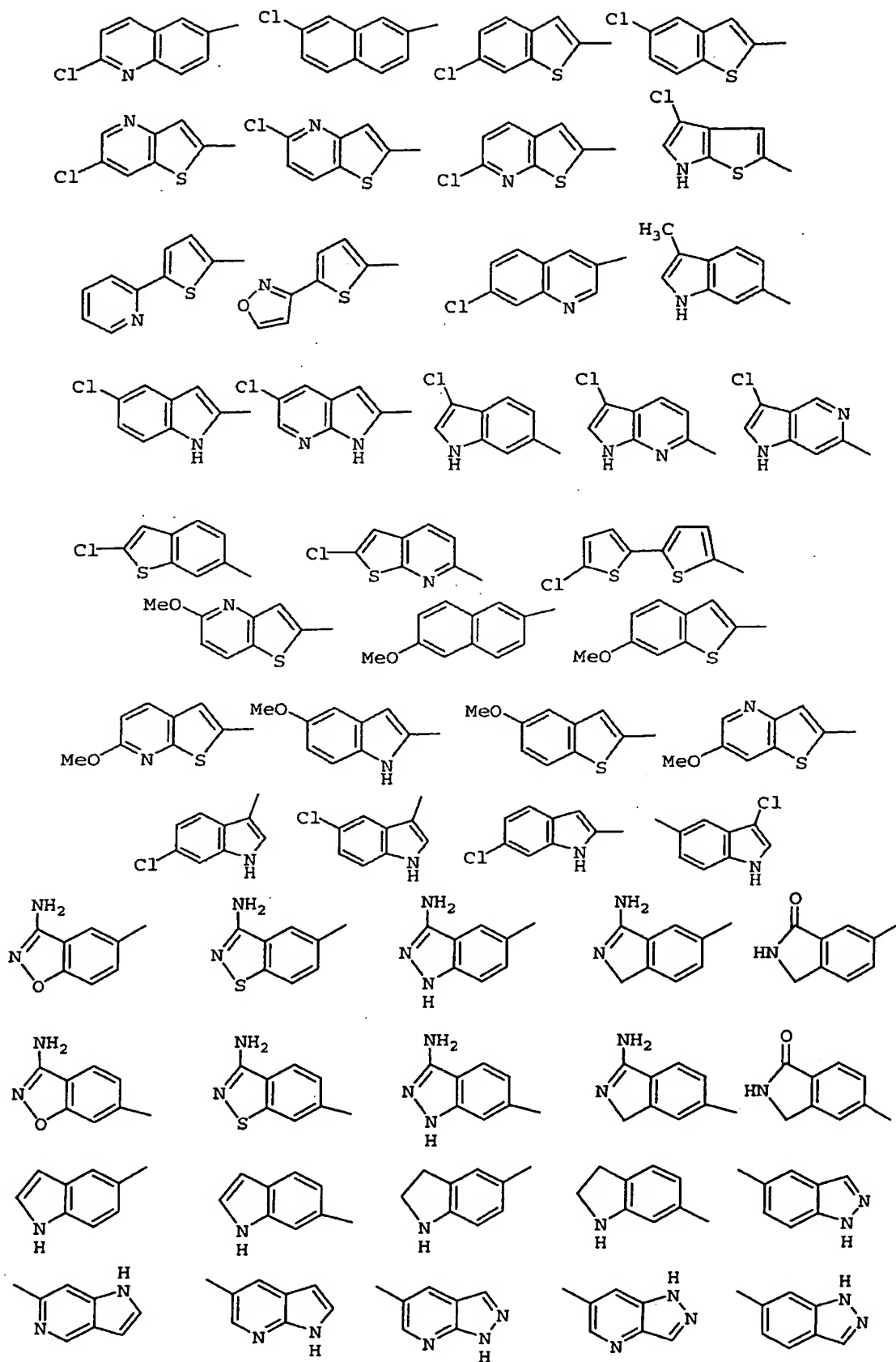
- phenyl; 2,5-bis-aminomethyl-phenyl;
- 5 2-amido-4-methoxy-phenyl; 2-amido-5-chloro-phenyl;
 - 2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
 - 2-aminomethyl-3-methoxy-phenyl;
 - 2-aminomethyl-4-fluoro-phenyl;
 - 2-aminomethyl-4-methoxy-phenyl;
 - 10 2-aminomethyl-5-fluoro-phenyl;
 - 2-aminomethyl-5-methoxy-phenyl;
 - 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 - 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
 - 2-aminosulfonyl-phenyl; 2-aminomethyl-4-ethyl-phenyl; 2-
 - 15 aminosulfonyl-4-ethyl-phenyl; 2-amido-4-ethyl-phenyl;
 - 2-hydroxy-4-methoxy-phenyl; 2-methylsulfonyl-phenyl;
 - 3-(N,N-dimethylamino)-4-chloro-phenyl;
 - 3-(N,N-dimethylamino)-phenyl; 3-(N-hydroxy-amidino)-phenyl;
 - 3-(N-methoxy-amidino)-phenyl;
 - 20 3-(N-methylamino)-4-chloro-phenyl;
 - 3-(N-methylamino)-phenyl; 3-amidino-phenyl;
 - 3-amido-6-hydroxy-phenyl; 3-amido-phenyl;
 - 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 - 3-amino-phenyl; 3-chloro-4-fluoro-phenyl; 3-chloro-phenyl;

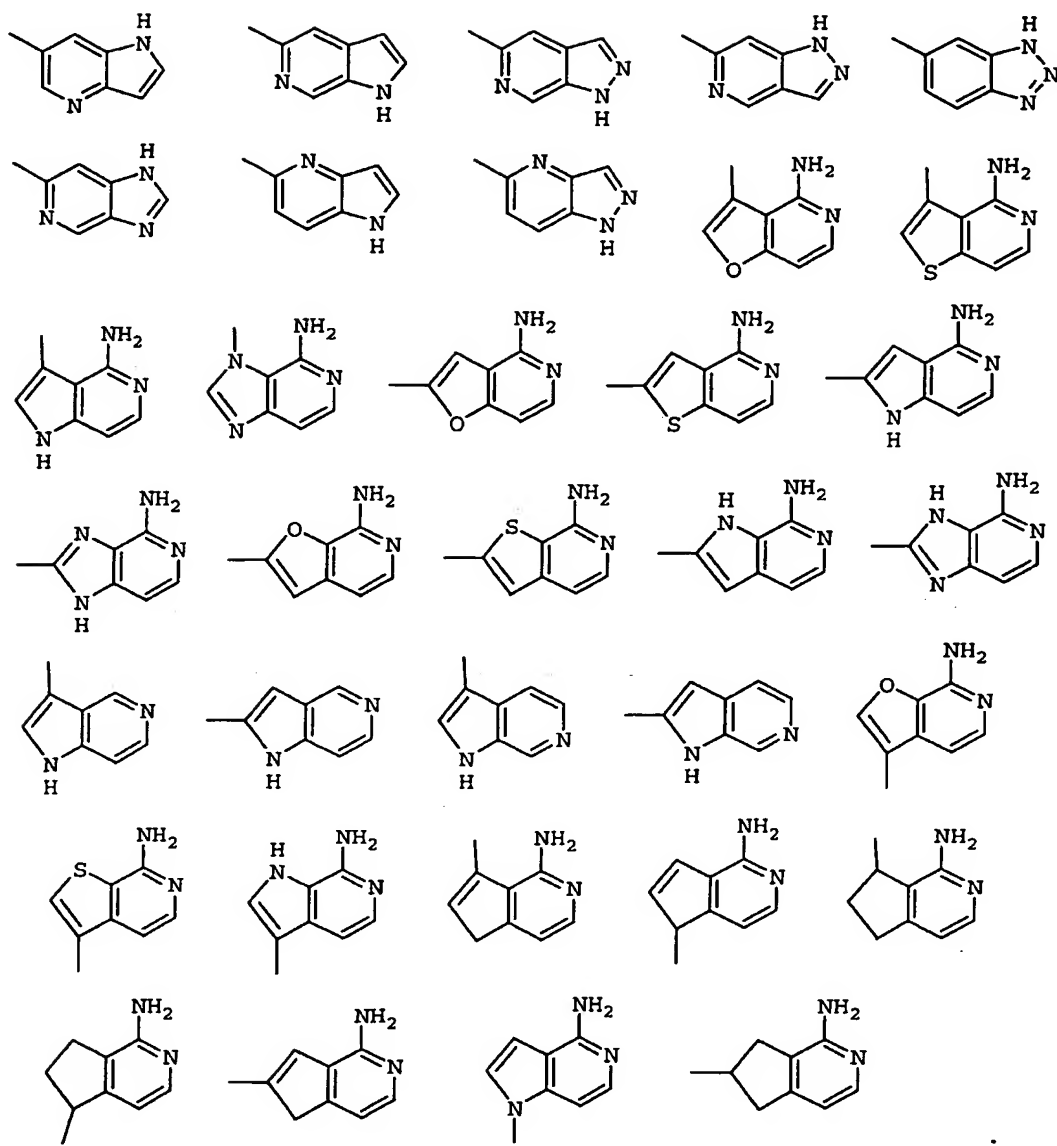
- 3-hydroxy-4-methoxy-phenyl;
 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
 4-(N-methylamino)-5-chloro-thien-2-yl;
 4-amino-5-chloro-thien-2-yl; 4-amino-pyrid-2-yl;
 5 4-chloro-3-fluoro-phenyl; 4-chloro-phenyl;
 4-chloro-pyrid-2-yl; 4-ethyl-phenyl; 4-ethyl-2-methylsulfonyl-phenyl;
 4-ethyl-2-methoxy-phenyl;
 4-methoxy-2-methylsulfonyl-phenyl; 4-methoxy-phenyl;
 2-methoxy-pyrid-5-yl;
 10 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
 5-(N-methylamino)-4-chloro-thien-2-yl;
 5-amino-4-chloro-thien-2-yl;
 5-chloro-2-aminosulfonyl-phenyl;
 5-chloro-2-methylsulfonyl-phenyl; 5-chloro-pyrid-2-yl;
 15 5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
 5-methyl-thien-2-yl; 5-fluoro-thien-2-yl;
 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;











- 5 G_1 is absent or is selected from $(CR^3R^3a)_{1-3}$, $CR^3=CR^3$,
 $(CR^3R^3a)_u C(O)(CR^3R^3a)_w$, $(CR^3R^3a)_u O(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^{3b}(CR^3R^3a)_w$, $(CR^3R^3a)_u C(O)NR^{3b}(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^{3bc}(O)(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^{3bc}(O)(CR^3R^3a)_u C(O)NR^{3b}(CR^3R^3a)_w$,
10 $(CR^3R^3a)_u S(CR^3R^3a)_w$, $(CR^3R^3a)_u S(O)(CR^3R^3a)_w$,
 $(CR^3R^3a)_u S(O)_2(CR^3R^3a)_w$, $(CR^3R^3a)_u S(O)NR^{3b}(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^{3bs}(O)_2(CR^3R^3a)_w$, and
 $(CR^3R^3a)_u S(O)_2NR^{3b}(CR^3R^3a)_w$, wherein $u + w$ total 0, 1,

or 2, provided that G₁ does not form a N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

5 A is selected from one of the following carbocycles and heterocycles which are substituted with 0-2 R⁴;

cyclohexyl, phenyl, piperidiny1, piperazinyl, pyridyl, pyrimidyl, furanyl, morpholinyl, thienyl, pyrrolyl, pyrrolidinyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyrazolyl, imidazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl, 1,3,4-triazolyl, benzofuranyl, benzothiofuranyl, indolinyl, indolyl, benzimidazolyl, benzoxazolyl, benzthiazolyl, indazolyl, benzisoxazolyl, benzisothiazolyl, and isoindazolyl;

20

X is selected from -(CR²R^{2a})₁₋₂-, -C(O)-, -S(O)₂-, -NR²S(O)₂-, -NR²S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR², -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

25 Y is a C₃₋₆ monocyclic carbocycle or 5-6 membered monocyclic heterocycle, wherein the carbocycle or heterocycle consists of carbon atoms and 0-2 heteroatoms selected from N, O, and S(O)_p, the carbocycle or heterocycle further comprises 0-1 double bonds and 0-1 carbonyl groups, and the carbocycle or heterocycle is substituted with 0-2 R⁴;

30

alternatively, Y is CY¹Y², and Y¹ and Y² are independently C₁₋₂ alkyl substituted with 0-1 R⁴;

35

R^{1a} , at each occurrence, is selected from H, R^{1b} , $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, CH_2R^{1b} , and $CH_2CH_2R^{1b}$, provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

5 alternatively, when two R^{1a} groups are attached to adjacent atoms or to the same carbon atom, together with the atoms to which they are attached they form a 5-6 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N,
10 O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b} and 0-3 ring double bonds;

R^{1b} is selected from H, CH_3 , CH_2CH_3 , F, Cl, Br, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} ,
15 $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, phenyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2
20 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

R^2 , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with
25 0-2 R^{4b} , benzyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

30 R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, phenyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon

atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom
5 to which they are attached, combine to form a 5 or 6
membered saturated, partially saturated or unsaturated
ring substituted with 0-2 R^{4b} and consisting of: 0-1
additional heteroatoms selected from the group
consisting of N, O, and S(O)_p;

10

R^{2b}, at each occurrence, is selected from CF₃, C₁₋₄ alkoxy,
CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl
substituted with 0-2 R^{4b}, and 5-6 membered aromatic
heterocycle substituted with 0-2 R^{4b} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the
group consisting of N, O, and S(O)_p;

20

R^{2c}, at each occurrence, is selected from CF₃, OH, OCH₃,
OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃,
CH(CH₃)₂, benzyl, phenyl substituted with 0-2 R^{4b}, and
5-6 membered aromatic heterocycle substituted with 0-2
R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
O, and S(O)_p;

25

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl
substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted
with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with
0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2
30 R^{4c} and consisting of: carbon atoms and 1-4
heteroatoms selected from the group consisting of N,
O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle
substituted with 0-2 R^{4c} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group

consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

5 R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4
 10 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms
 15 other than a C(O)-halo or C(O)-S(O)_p moiety;

R⁴, at each occurrence, is selected from H, (CH₂)₂OR², CH₂OR², OR², F, Cl, Br, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃,
 20 C(CH₃)₃, -CN, NO₂, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, C(O)R^{2c}, NR²C(O)R^{2b}, C(O)NR²R^{2a}, SO₂NR²R^{2a}, CF₃, and CF₂CF₃;

R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle
 25 substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d}, (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d},
 30 (CR³R^{3g})_r-NR^{2d}C(O)R^{2e}, (CR³R^{3g})_r-C(O)R^{2e}, (CR³R^{3g})_r-OC(O)R^{2e}, (CR³R^{3g})_r-C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-C(O)OR^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)OR^{2d}, (CR³R^{3g})_r-SO₂NR^{2d}R^{2d},

$(\text{CR}^3\text{R}^3\text{g})_r\text{-NR}^{2d}\text{SO}_2\text{R}^{2d}$, and $(\text{CR}^3\text{R}^3\text{g})_r\text{-S(O)}_p\text{R}^{2d}$, provided that $\text{S(O)}_p\text{R}^{2d}$ forms other than $\text{S(O)}_2\text{H}$ or S(O)H ;

R^{4b} , at each occurrence, is selected from H, =O, OR^3 ,
 5 CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, -CN,
 NO_2 , NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, C(O)R^3 , $\text{CH}_2\text{-C(O)R}^3$, C(O)OR^{3c} ,
 $\text{CH}_2\text{-C(O)OR}^{3c}$, $\text{NR}^3\text{C(O)R}^{3a}$, $\text{CH}_2\text{NR}^3\text{C(O)R}^{3a}$, $\text{C(O)NR}^3\text{R}^{3a}$, $\text{CH}_2\text{-}$
 $\text{C(O)NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{CH}_2\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl,
 $\text{CH}_2\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl, $\text{NR}^3\text{SO}_2\text{-phenyl}$, $\text{CH}_2\text{NR}^3\text{SO}_2\text{-phenyl}$,
 10 $\text{S(O)}_p\text{CF}_3$, $\text{CH}_2\text{S(O)}_p\text{CF}_3$, $\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $\text{CH}_2\text{S(O)}_p\text{-C}_{1-4}$
 alkyl, $\text{S(O)}_p\text{-phenyl}$, $\text{CH}_2\text{S(O)}_p\text{-phenyl}$, and CF_3 ;

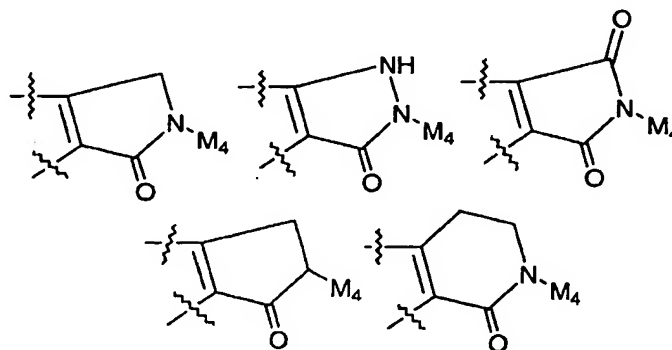
R^{4c} , at each occurrence, is selected from =O, OR^2 ,
 $(\text{CR}^3\text{R}^{3a})\text{OR}^2$, F, $(\text{CR}^3\text{R}^{3a})\text{F}$, Br, $(\text{CR}^3\text{R}^{3a})\text{Br}$, Cl,
 15 $(\text{CR}^3\text{R}^{3a})\text{Cl}$, CF_3 , $(\text{CR}^3\text{R}^{3a})\text{CF}_3$, C_{1-4} alkyl, C_{2-3} alkenyl,
 C_{2-3} alkynyl, -CN, $(\text{CR}^3\text{R}^{3a})\text{CN}$, NO_2 , $(\text{CR}^3\text{R}^{3a})\text{NO}_2$, NR^2R^{2a} ,
 $(\text{CR}^3\text{R}^{3a})\text{NR}^2\text{R}^{2a}$, $\text{N}(\rightarrow\text{O})\text{R}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{N}(\rightarrow\text{O})\text{R}^2\text{R}^{2a}$, C(O)R^{2c} ,
 $(\text{CR}^3\text{R}^{3a})\text{C(O)R}^{2c}$, $\text{NR}^2\text{C(O)R}^{2b}$, $(\text{CR}^3\text{R}^{3a})\text{NR}^2\text{C(O)R}^{2b}$,
 $\text{C(O)NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{C(O)NR}^2\text{R}^{2a}$, $\text{NR}^2\text{C(O)NR}^2\text{R}^{2a}$,
 20 $(\text{CR}^3\text{R}^{3a})\text{NR}^2\text{C(O)NR}^2\text{R}^{2a}$, $\text{SO}_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{SO}_2\text{NR}^2\text{R}^{2a}$,
 $\text{NR}^2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{NR}^2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{NR}^2\text{SO}_2\text{R}^{5a}$,
 $(\text{CR}^3\text{R}^{3a})\text{NR}^2\text{SO}_2\text{R}^{5a}$, $\text{S(O)}_p\text{R}^{5a}$, $(\text{CR}^3\text{R}^{3a})\text{S(O)}_p\text{R}^{5a}$, CF_3 ,
 CF_2CF_3 , C_{3-10} carbocycle substituted with 0-2 R^{4b} ,
 $(\text{CR}^3\text{R}^{3a})\text{-C}_{3-10}$ carbocycle substituted with 0-2 R^{4b} , 5-10
 25 membered heterocycle substituted with 0-2 R^{4b} and
 consisting of carbon atoms and from 1-4 heteroatoms
 selected from the group consisting of N, O, and S(O)_p ,
 and $(\text{CR}^3\text{R}^{3a})\text{-5-10}$ membered heterocycle substituted with
 0-2 R^{4b} and consisting of carbon atoms and from 1-4
 30 heteroatoms selected from the group consisting of N,
 O, and S(O)_p ;

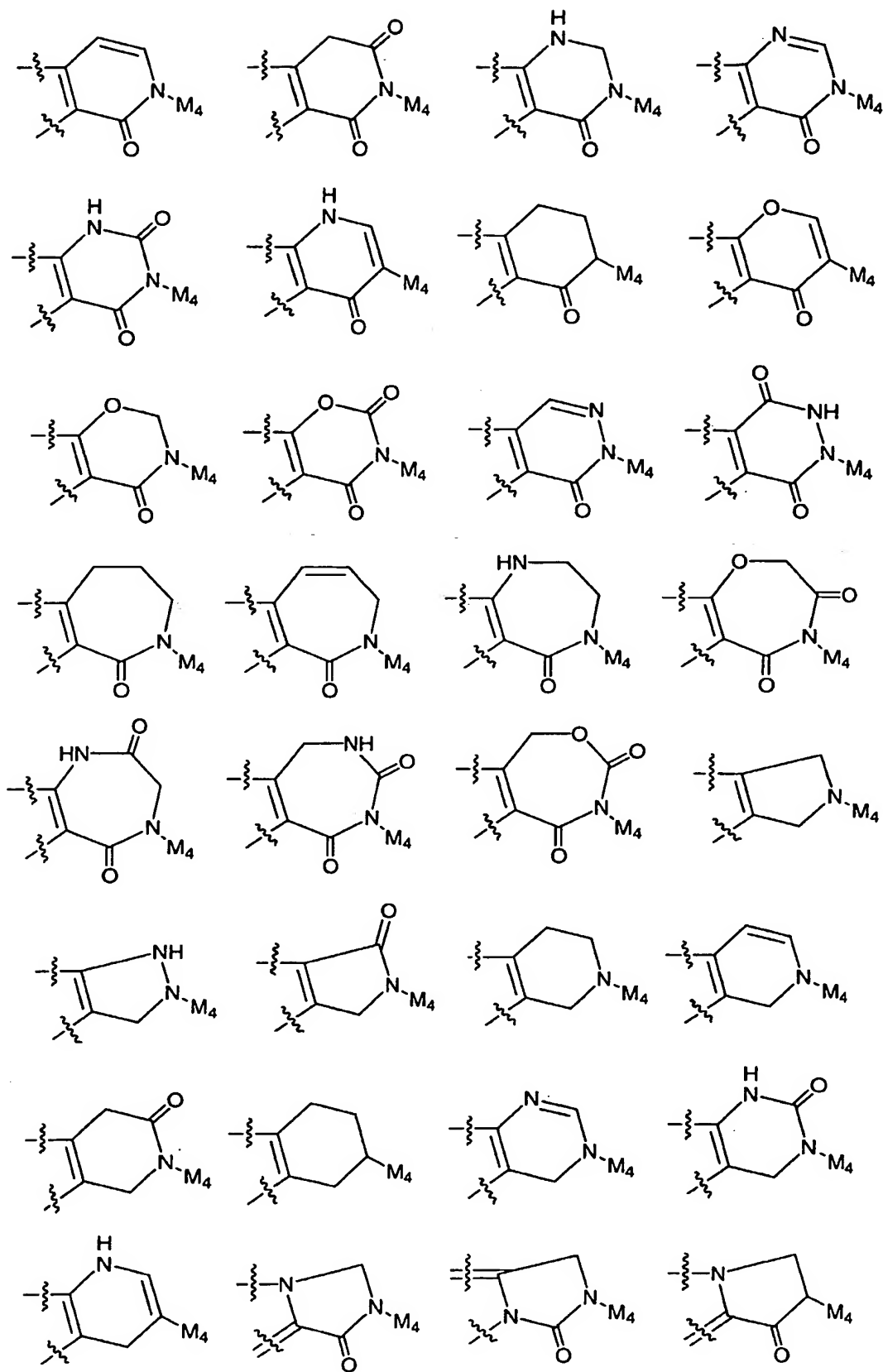
- R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$, $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 5 $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -phenyl, $S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 , phenyl substituted with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and benzyl substituted with 0-2 R^6 ; and,
- 10 R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$, $SO_2NR^2R^{2a}$, and $NR^2SO_2C_{1-4}$ alkyl.

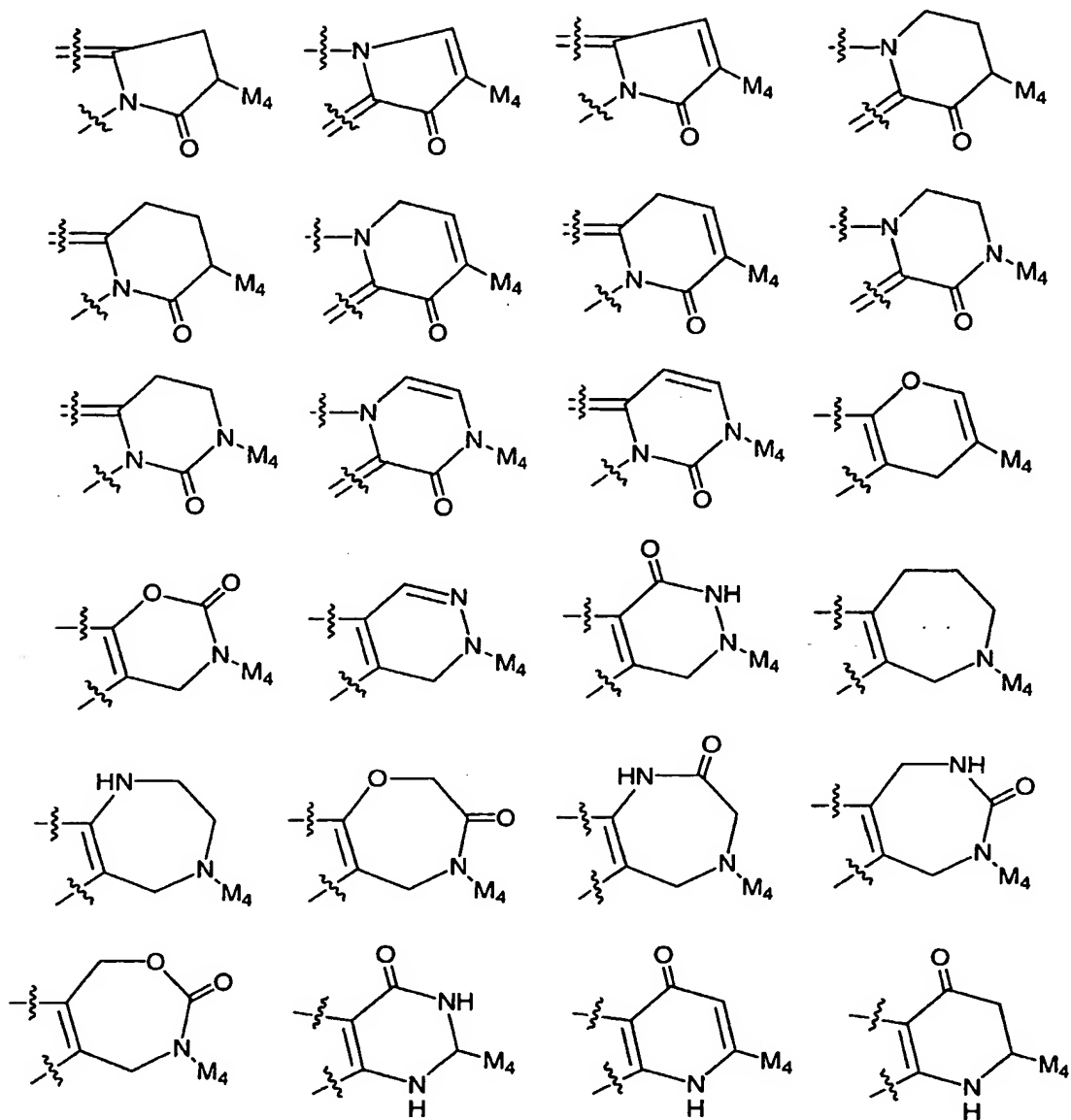
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[4] In another preferred embodiment, the present invention provides a novel compound, wherein:

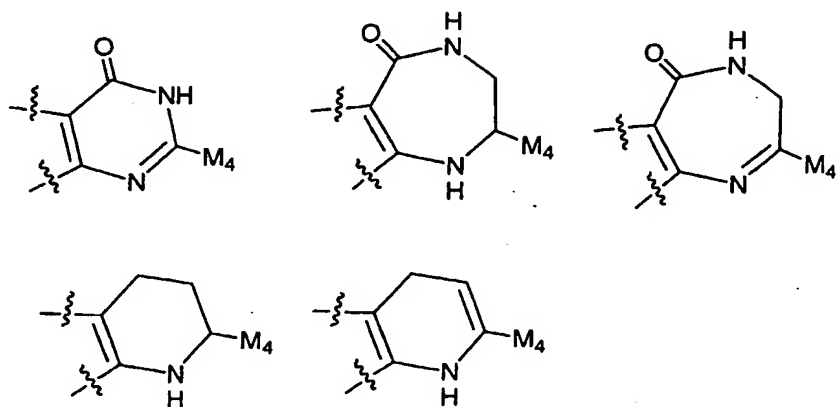
20 ring M is substituted with 0-2 R^{1a} and is selected from the group:



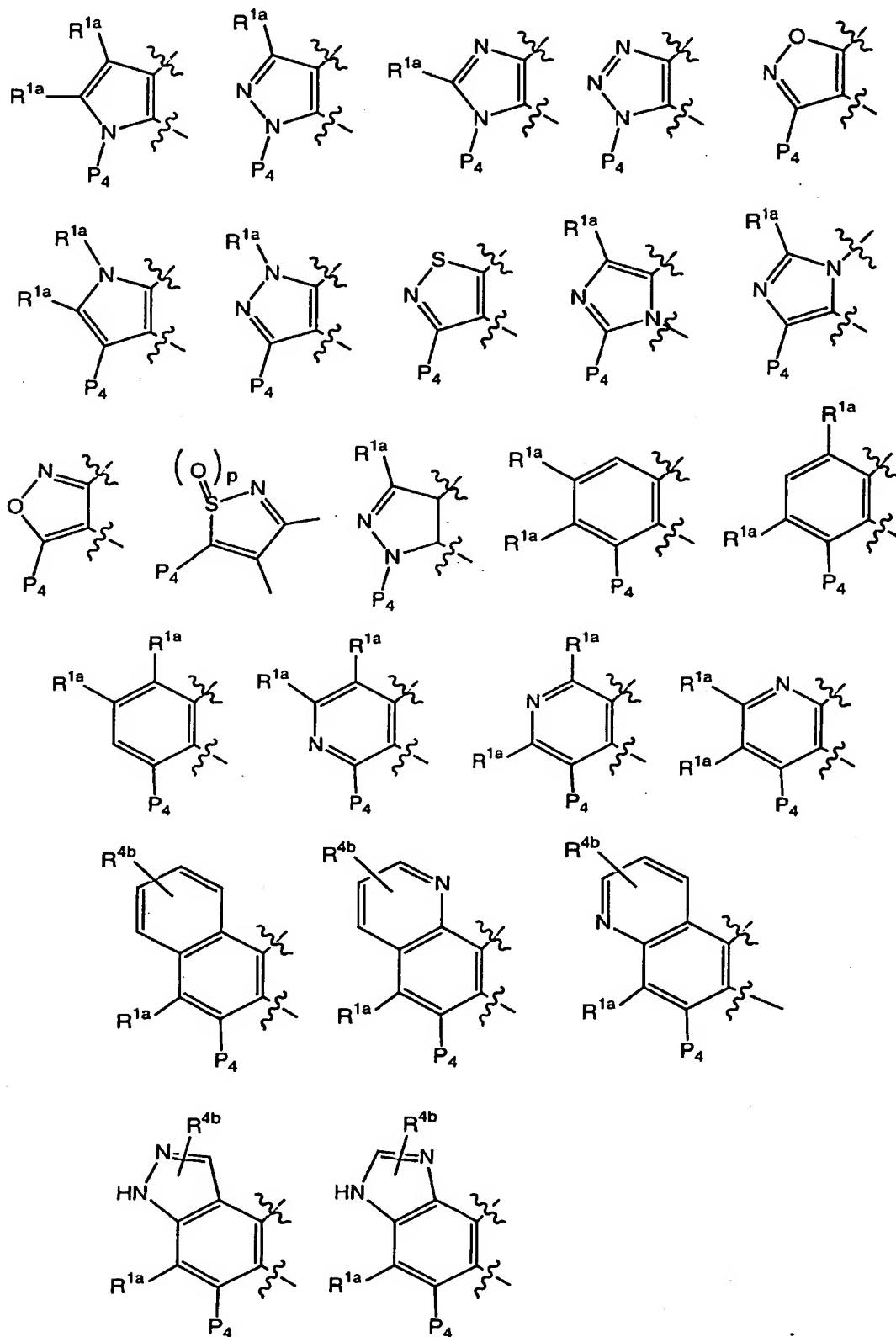




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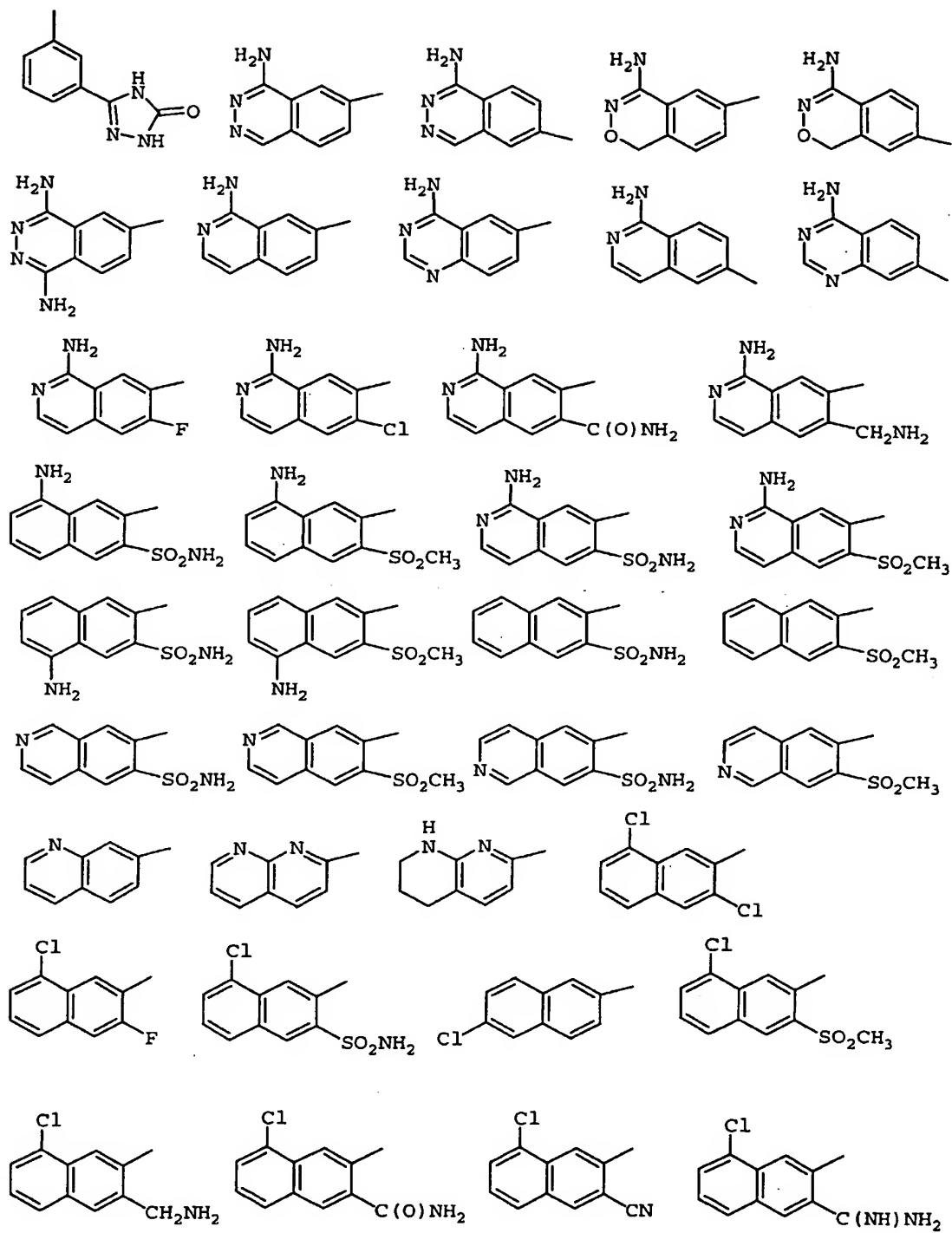
ring P, including P₁, P₂, P₃, and P₄ is selected from group:

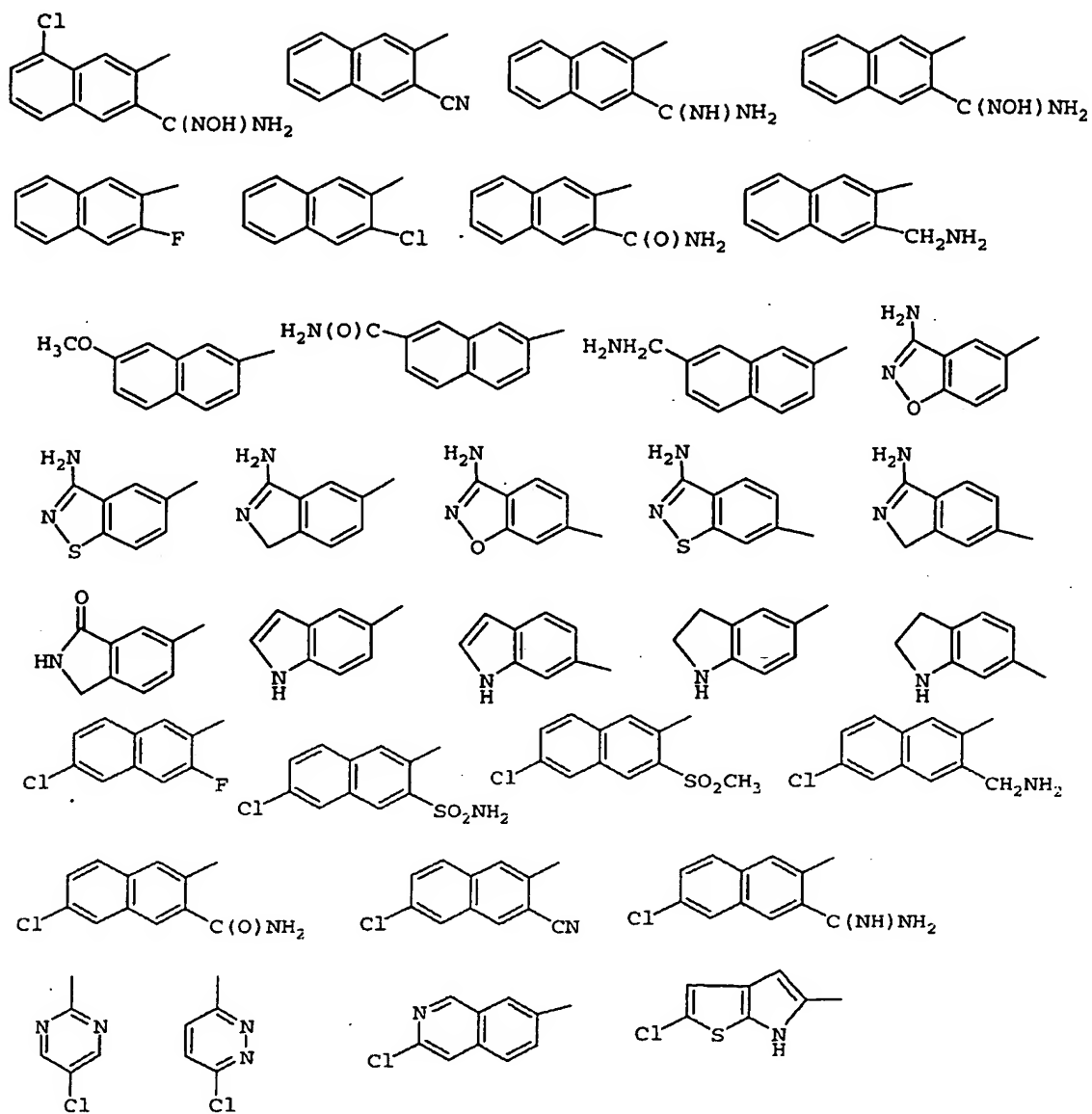


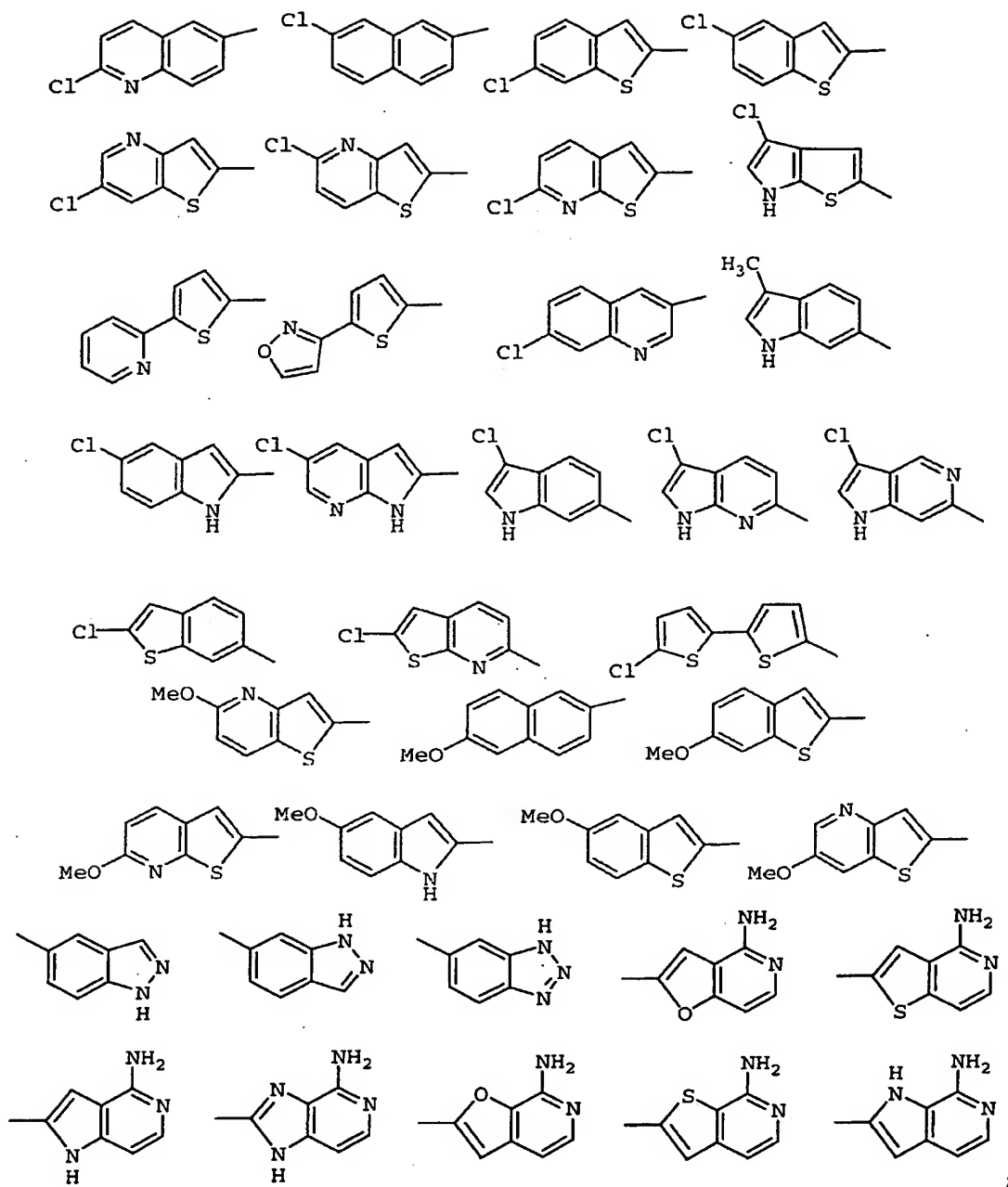
;

G is selected from the group:

- phenyl; 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
- 2-aminomethyl-3-fluoro-phenyl;
- 5 2-aminomethyl-4-fluoro-phenyl;
- 2-aminomethyl-4-methoxy-phenyl;
- 2-aminomethyl-5-fluoro-phenyl;
- 2-aminomethyl-5-methoxy-phenyl;
- 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
- 10 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
- 2-aminosulfonyl-phenyl; 2-methylsulfonyl-phenyl; 2-
- aminomethyl-4-ethyl-phenyl; 2-aminosulfonyl-4-ethyl-phenyl;
- 2-amido-4-ethyl-phenyl;
- 3-(N,N-dimethylamino)-4-chloro-phenyl;
- 15 3-(N,N-dimethylamino)-phenyl;
- 3-(N-methylamino)-4-chloro-phenyl;
- 3-(N-methylamino)-phenyl; 3-amido-phenyl;
- 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
- 3-amino-phenyl; 3-chloro-phenyl;
- 20 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
- 4-(N-methylamino)-5-chloro-thien-2-yl;
- 4-amino-5-chloro-thien-2-yl; 4-chloro-phenyl; 4-ethyl-
- phenyl; 4-ethyl-2-methylsulfonyl-phenyl; 4-ethyl-2-methoxy-
- phenyl; 4-methoxy-2-methylsulfonyl-phenyl;
- 25 4-methoxy-phenyl;
- 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
- 5-(N-methylamino)-4-chloro-thien-2-yl;
- 5-amino-4-chloro-thien-2-yl; 5-chloro-pyrid-2-yl;
- 5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
- 30 5-methyl-thien-2-yl; 5-fluoro-thien-2-yl;
- 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;







- 5 G_1 is absent or is selected from CH_2 , CH_2CH_2 , $CH=CH$, CH_2O , OCH_2 , NH , CH_2NH , $NHCH_2$, $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $CH_2S(O)_2$, $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$, provided that G_1 does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;

- A is selected from cyclohexyl, piperidinyl, phenyl, pyridyl, and pyrimidyl, and is substituted with 0-2 R^4 ;
- 5 X is selected from CH_2 , $C(O)$, $-S(O)_2-$, $-NHC(O)-$, $-C(O)NH-$, $-CH_2NH-$, O, and $-CH_2O-$;
- Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentanonyl, cyclohexyl, cyclohexanonyl, pyrrolidinyl, pyrrolidinonyl, 10 piperidinyl, piperidinonyl, tetrahydrofuranyl, and tetrahydropyranyl, and, when Y is a ring, Y is substituted with 0-1 R^4 ;
- 15 R^{1a} , at each occurrence, is selected from H, R^{1b} , $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, and CH_2R^{1b} , provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;
- R^{1b} is selected from CH_3 , CH_2CH_3 , F, Cl, Br, $-CN$, CF_3 , OR^2 , 20 NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , CO_2R^{2a} , $S(O)_pR^2$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided 25 that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;
- R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-1 R^{4b} , 30 benzyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2a}, at each occurrence, is selected from H, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2b}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2c}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4

heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

R⁴, at each occurrence, is selected from OH, OR², CH₂OR², (CH₂)₂OR², F, Br, Cl, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, CF₃, and CF₂CF₃;

R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d}, (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d}, (CR³R^{3g})_r-C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)R^{2e},

$(\text{CR}^3\text{R}^3\text{g})_x\text{-C(O)R}^{2e}$, $(\text{CR}^3\text{R}^3\text{g})_x\text{-NR}^{2d}\text{C(O)NR}^{2d}\text{R}^{2d}$,
 $(\text{CR}^3\text{R}^3\text{g})_x\text{-NR}^{2d}\text{C(O)OR}^{2d}$, $(\text{CR}^3\text{R}^3\text{g})_x\text{-NR}^{2d}\text{SO}_2\text{R}^{2d}$, and
 $(\text{CR}^3\text{R}^3\text{g})_x\text{-S(O)}_p\text{R}^{2d}$, provided that $\text{S(O)}_p\text{R}^{2d}$ forms other
than $\text{S(O)}_2\text{H}$ or S(O)H ;

5

R^{4b} , at each occurrence, is selected from H, =O, OR^3 ,
 CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, -CN,
 NO_2 , NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, C(O)R^3 , C(O)OR^{3c} , $\text{NR}^3\text{C(O)R}^{3a}$,
 $\text{C(O)NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl, $\text{NR}^3\text{SO}_2\text{-phenyl}$,
10 $\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $\text{S(O)}_p\text{-phenyl}$, and CF_3 ;

R^{4c} , at each occurrence, is selected from =O, OR^2 , CH_2OR^2 ,
F, Br, Cl, CF_3 , CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, C_{2-3}
alkenyl, C_{2-3} alkynyl, -CN, NO_2 , NR^2R^{2a} , $\text{CH}_2\text{NR}^2\text{R}^{2a}$,
15 $\text{N}(\rightarrow\text{O})\text{R}^2\text{R}^{2a}$, $\text{CH}_2\text{N}(\rightarrow\text{O})\text{R}^2\text{R}^{2a}$, C(O)R^{2c} , $\text{CH}_2\text{C(O)R}^{2c}$,
 $\text{NR}^2\text{C(O)R}^{2b}$, $\text{CH}_2\text{NR}^2\text{C(O)R}^{2b}$, $\text{C(O)NR}^2\text{R}^{2a}$, $\text{CH}_2\text{C(O)NR}^2\text{R}^{2a}$,
 $\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{CH}_2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{NR}^2\text{SO}_2\text{R}^{5a}$, $\text{CH}_2\text{NR}^2\text{SO}_2\text{R}^{5a}$,
 $\text{S(O)}_p\text{R}^{5a}$, $\text{CH}_2\text{S(O)}_p\text{R}^{5a}$, CF_3 , CF_2CF_3 , C_{3-6} carbocycle
substituted with 0-2 R^{4b} , $(\text{CH}_2)\text{-C}_{3-6}$ carbocycle
20 substituted with 0-2 R^{4b} , 5-6 membered heterocycle
substituted with 0-2 R^{4b} and consisting of carbon atoms
and from 1-4 heteroatoms selected from the group
consisting of N, O, and S(O)_p , and $(\text{CH}_2)\text{-5-6}$ membered
heterocycle substituted with 0-2 R^{4b} and consisting of
25 carbon atoms and from 1-4 heteroatoms selected from
the group consisting of N, O, and S(O)_p ;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
30 NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, C(O)R^3 , C(O)OR^{3c} , $\text{NR}^3\text{C(O)R}^{3a}$,
 $\text{C(O)NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl, $\text{NR}^3\text{SO}_2\text{-phenyl}$,
 $\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $\text{S(O)}_p\text{-phenyl}$, CF_3 , phenyl substituted

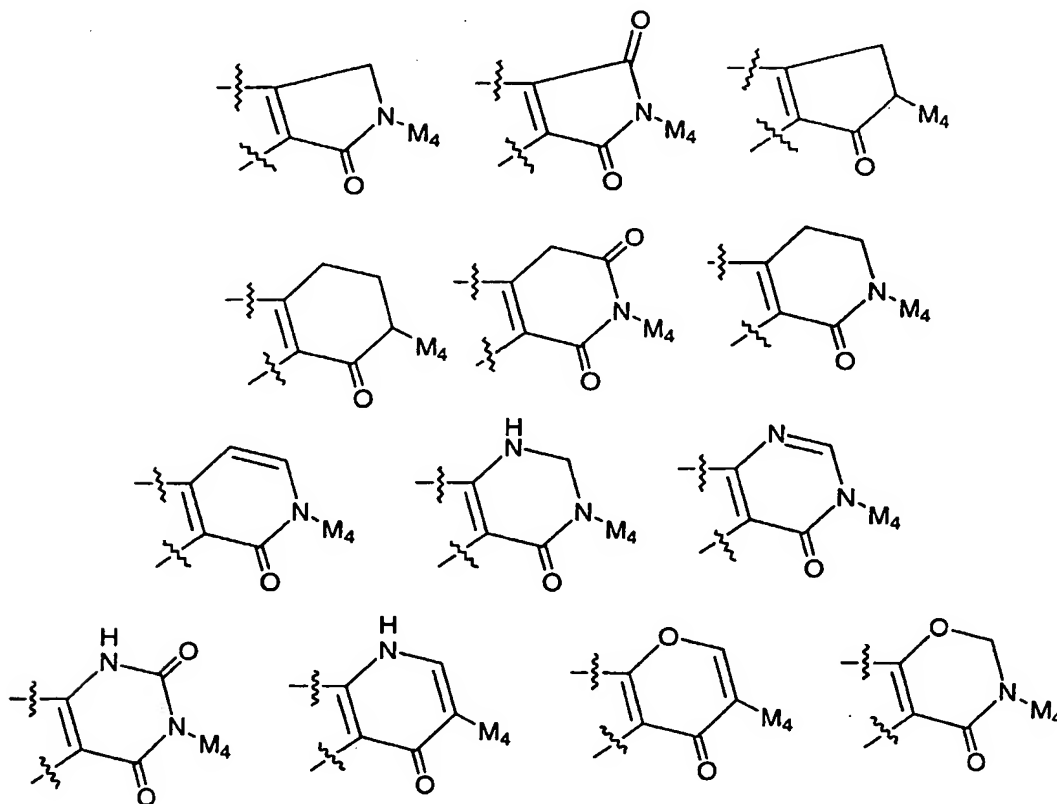
with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
benzyl substituted with 0-2 R^6 ;

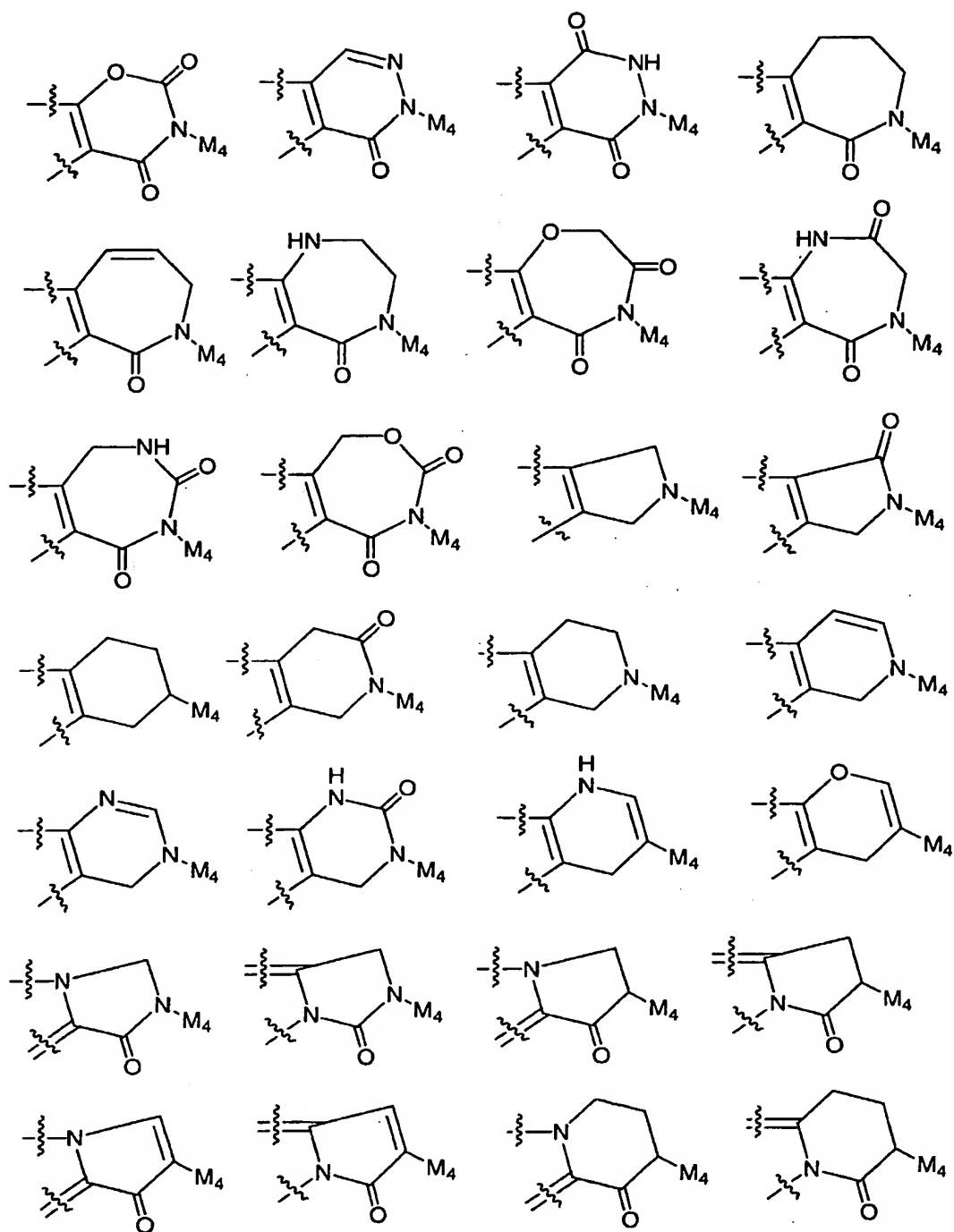
5 R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $-CN$, NO_2 , NR^2R^{2a} ,
 $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$, and
 $SO_2NR^2R^{2a}$; and,

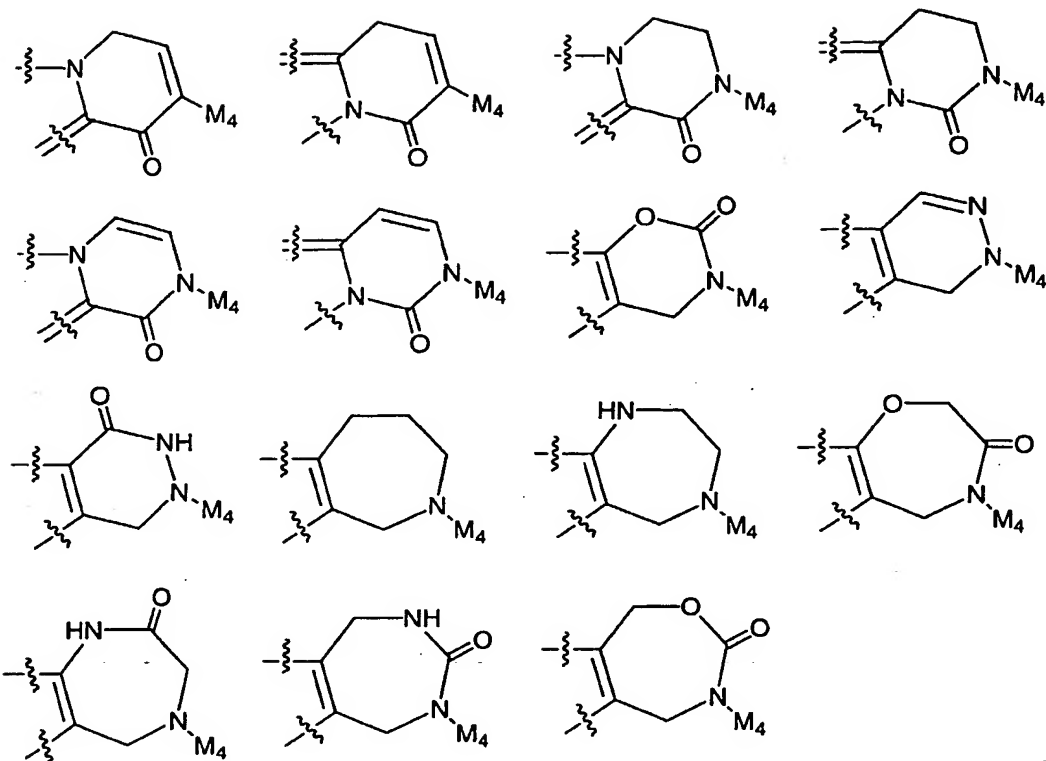
10 r, at each occurrence, is selected from 0, 1, and 2.

[5] In another preferred embodiment, the present invention
provides a novel compound, wherein:

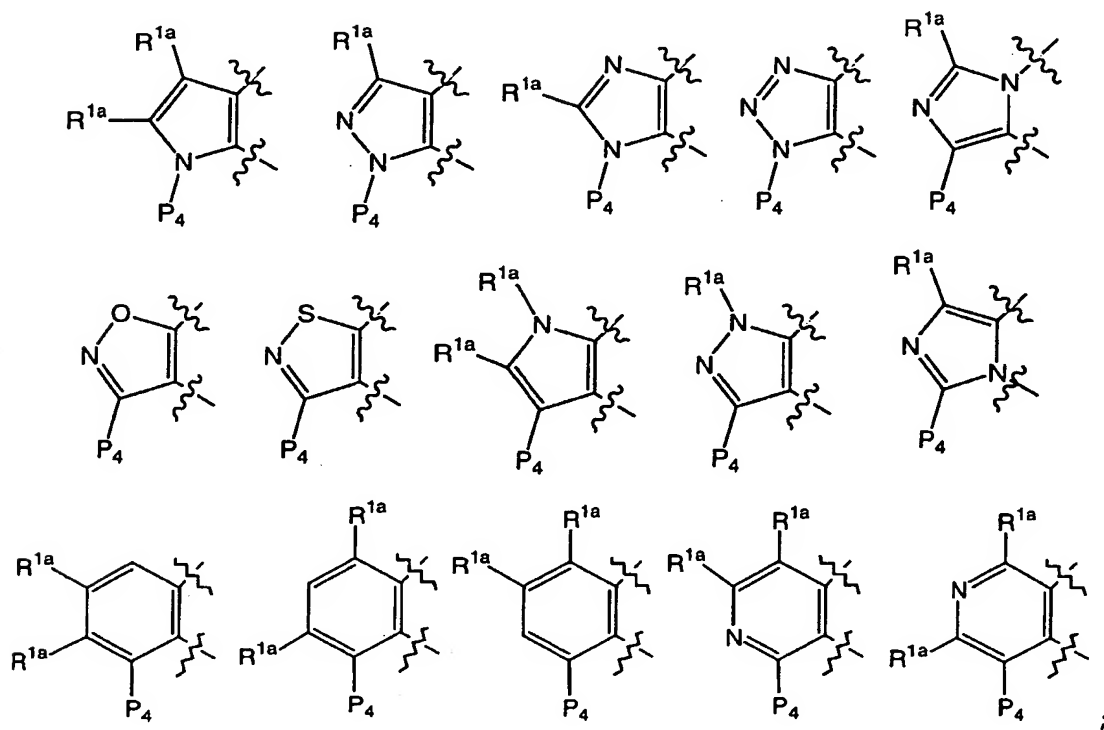
15 ring M is substituted with 0-1 R^{1a} and is selected from the
group:





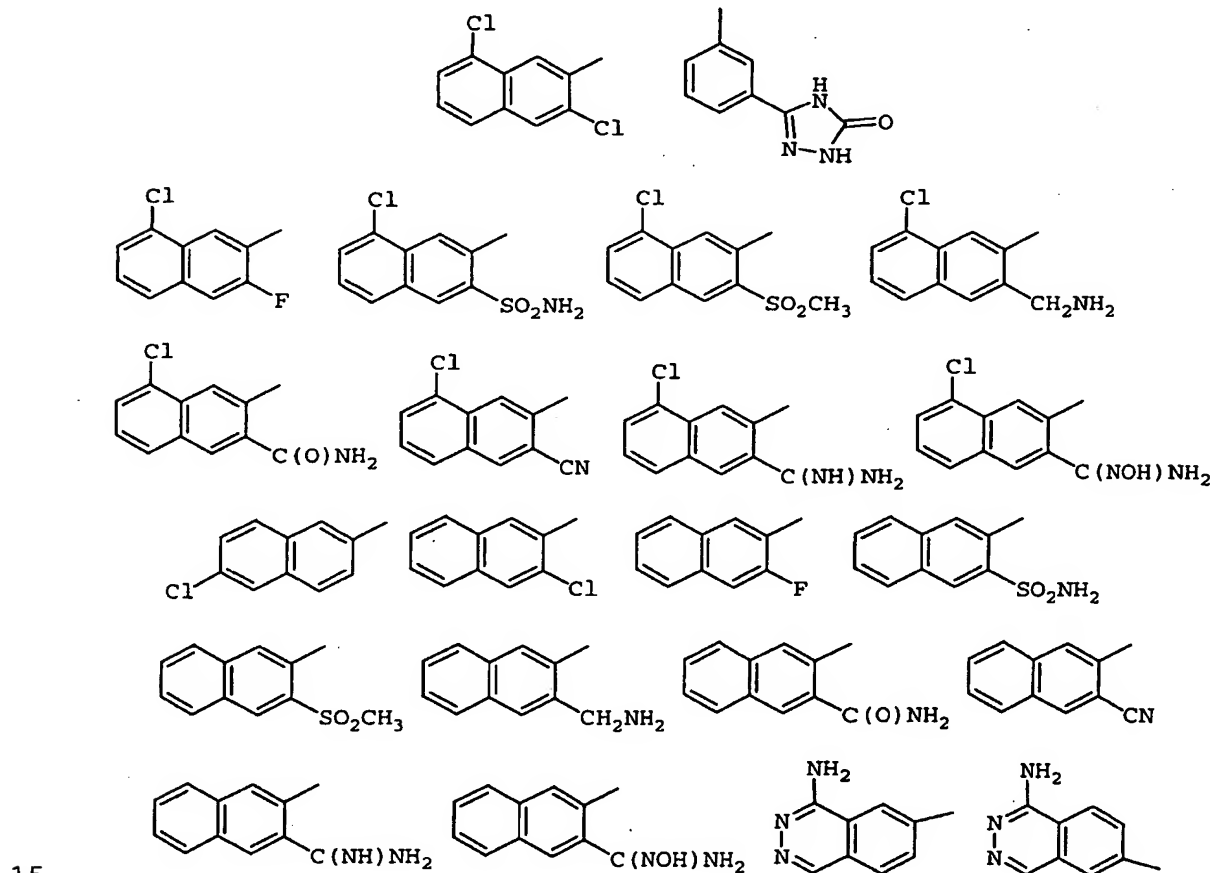


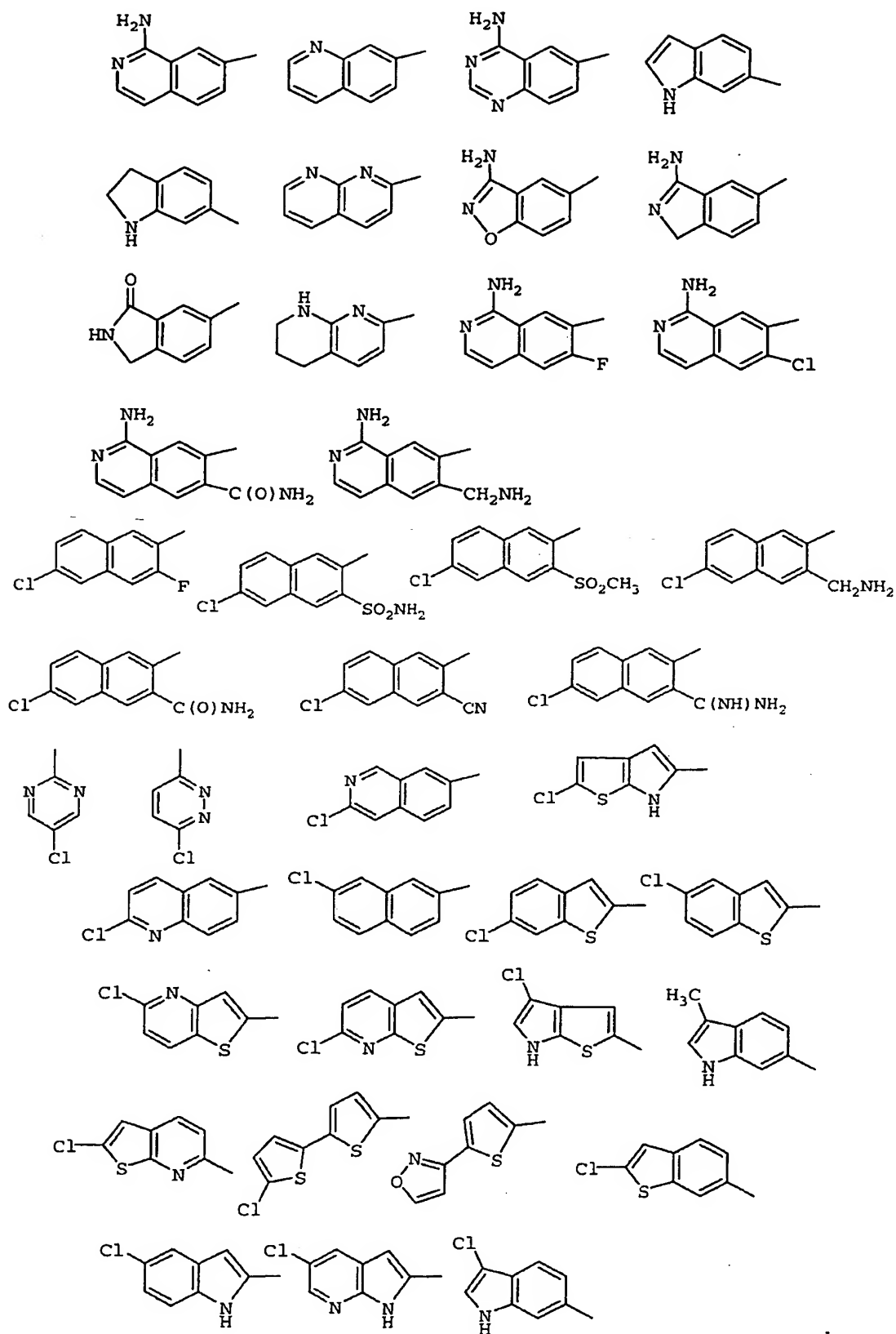
ring P, including P₁, P₂, P₃, and P₄ is selected from group:



-G is selected from:

- 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
 2-aminomethyl-3-fluoro-phenyl;
 2-aminomethyl-4-fluoro-phenyl;
 5 2-aminomethyl-5-fluoro-phenyl;
 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
 2-aminosulfonyl-phenyl; 3-amido-phenyl;
 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 10 3-chloro-phenyl; 4-chloro-phenyl; 4-ethyl-phenyl;
 4-methoxy-phenyl; 5-chloro-pyrid-2-yl; 5-chloro-thien-2-yl;
 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;





A is selected from the group: cyclohexyl, piperidiny, phenyl, 2-pyridyl, 3-pyridyl, 2-pyrimidyl, 2-Cl-phenyl, 3-Cl-phenyl, 2-F-phenyl, 3-F-phenyl, 2-methylphenyl, 2-aminophenyl, and 2-methoxyphenyl;

5

Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, 2-cyclopentanonyl, cyclohexyl, 2-cyclohexanonyl, pyrrolidiny (attached to A and R^{4a} at the 2-position), pyrrolidiny (attached to A and R^{4a} at the 3-position), 2-pyrrolidinonyl (attached to A and R^{4a} at the 3-position), piperidiny (attached to A and R^{4a} at the 4-position), 4-piperidinonyl (attached to A and R^{4a} at the 3-position), tetrahydrofuranyl, and tetrahydropyranyl (attached to A and R^{4a} at the 4-position);

10

15

R^{1a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, CH_2F , CH_2Cl , Br, CH_2Br , -CN, CH_2CN , CF_3 , CH_2CF_3 , OCH_3 , CH_2OH , $C(CH_3)_2OH$, CH_2OCH_3 , NH_2 , CH_2NH_2 , $NHCH_3$, CH_2NHCH_3 , $N(CH_3)_2$, $CH_2N(CH_3)_2$, CO_2H , $COCH_3$, CO_2CH_3 , $CH_2CO_2CH_3$, SCH_3 , CH_2SCH_3 , $S(O)CH_3$, $CH_2S(O)CH_3$, $S(O)_2CH_3$, $CH_2S(O)_2CH_3$, $C(O)NH_2$, $CH_2C(O)NH_2$, SO_2NH_2 , $CH_2SO_2NH_2$, $NHSO_2CH_3$, $CH_2NHSO_2CH_3$, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyridin-2-yl-N-oxide, pyridin-3-yl-N-oxide, pyridin-4-yl-N-oxide, imidazol-1-yl, CH_2 -imidazol-1-yl, 4-methyl-oxazol-2-yl, 4-N,N-dimethylaminomethyl-oxazol-2-yl, 1,2,3,4-tetrazol-1-yl, 1,2,3,4-tetrazol-5-yl, CH_2 -1,2,3,4-tetrazol-1-yl, and CH_2 -1,2,3,4-tetrazol-5-yl, provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

20

25

30

R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-1 R^{4b} , benzyl substituted with 0-1 R^{4b} , and 5 membered

aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

- 5 R^{2a} , at each occurrence, is selected from H, CH_3 , and CH_2CH_3 ;

alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6
10 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

- 15 R^{2b} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , CH_3 , and CH_2CH_3 ;

R^{2c} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , CH_3 , and CH_2CH_3 ;

20

- R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c}
25 and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p$ - $S(O)_p$, S-O, O-N, O-S, or O-O moiety;

30

- R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c}

and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

5

R^{4a} is selected from -(CH₂)_r-5-6 membered carbocycle substituted with 0-3 R^{4c}, -(CH₂)_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CH₂)_rNR^{2d}R^{2d},
 10 (CH₂)_rN(→O)R^{2d}R^{2d}, (CH₂)_rOR^{2d}, (CH₂)_r-C(O)NR^{2d}R^{2d},
 (CH₂)_r-NR^{2d}C(O)R^{2e}, (CH₂)_r-C(O)R^{2e},
 (CH₂)_r-NR^{2d}C(O)NR^{2d}R^{2d}, (CH₂)_r-NR^{2d}C(O)OR^{2d},
 (CH₂)_r-NR^{2d}SO₂R^{2d}, and (CH₂)_r-S(O)_pR^{2d}, provided that
 15 S(O)_pR^{2d} forms other than S(O)₂H or S(O)H;

R^{4b}, at each occurrence, is selected from H, =O, OR³, CH₂OR³, F, Cl, CH₃, CH₂CH₃, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³,
 C(O)OR^{3c}, NR³C(O)R^{3a}, C(O)NR³R^{3a}, SO₂NR³R^{3a},
 20 NR³SO₂-phenyl, S(O)₂CH₃, S(O)₂-phenyl, and CF₃;

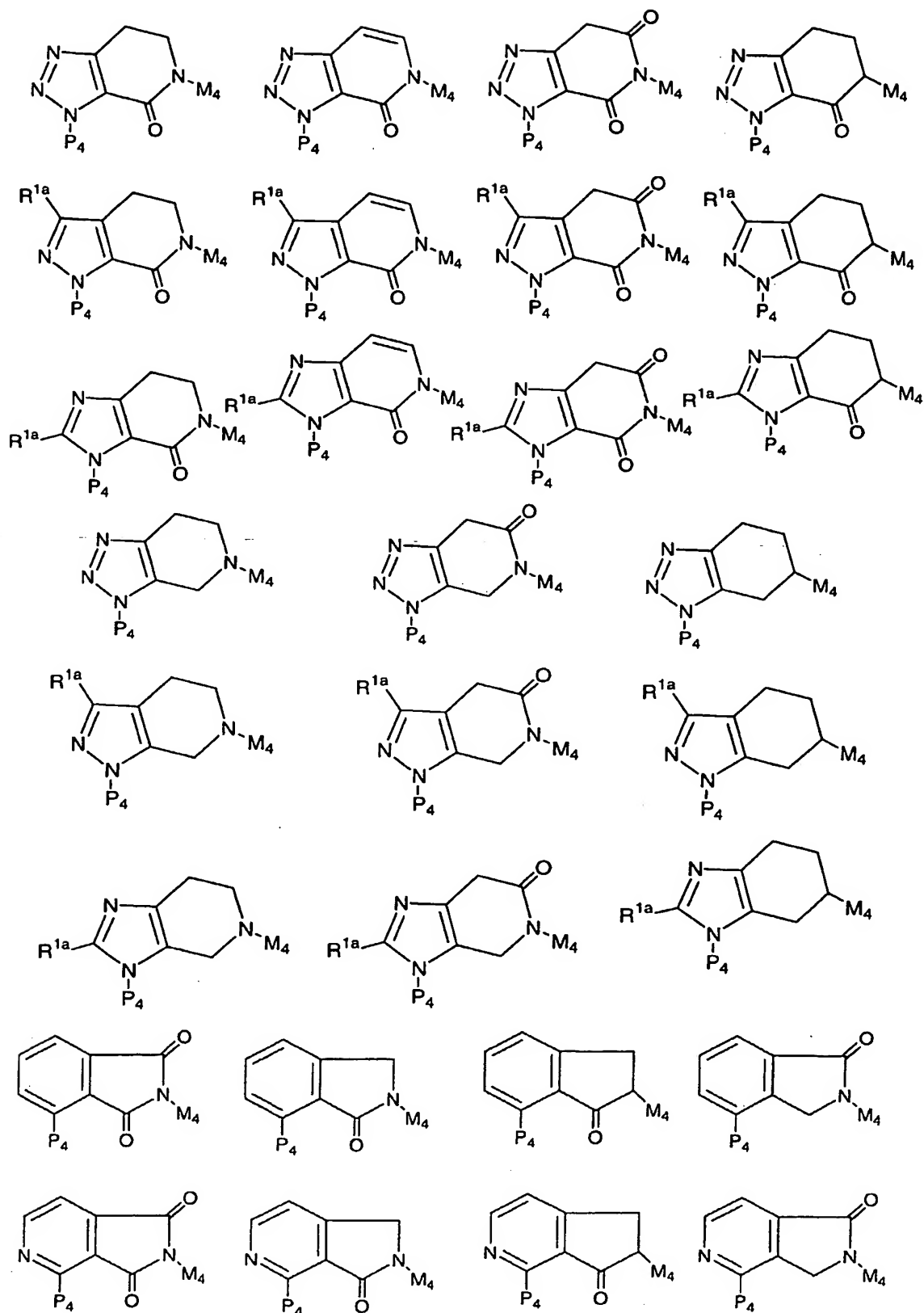
R^{4c}, at each occurrence, is selected from =O, OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, C₂₋₃ alkenyl, C₂₋₃ alkynyl, CH₂OH, CH₂OCH₃,
 25 CH₂OCH₂CH₃, CH₂OCH₂CH₂CH₃, CH₂OCH(CH₃)₂, F, Br, Cl, CF₃,
 NR²R^{2a}, CH₂NR²R^{2a}, N(→O)R²R^{2a}, CH₂N(→O)R²R^{2a}, C(O)R^{2c},
 CH₂C(O)R^{2c}, NR²C(O)R^{2b}, CH₂NR²C(O)R^{2b}, C(O)NR²R^{2a},
 CH₂C(O)NR²R^{2a}, SO₂NR²R^{2a}, CH₂SO₂NR²R^{2a}, NR²SO₂R^{5a},
 CH₂NR²SO₂R^{5a}, S(O)_pR^{5a}, CH₂S(O)_pR^{5a}, CF₃, cyclopropyl
 30 substituted with 0-1 R^{4b}, cyclobutyl substituted with 0-1 R^{4b}, cyclopentyl substituted with 0-1 R^{4b}, phenyl substituted with 0-1 R^{4b}, -CH₂-cyclopropyl substituted

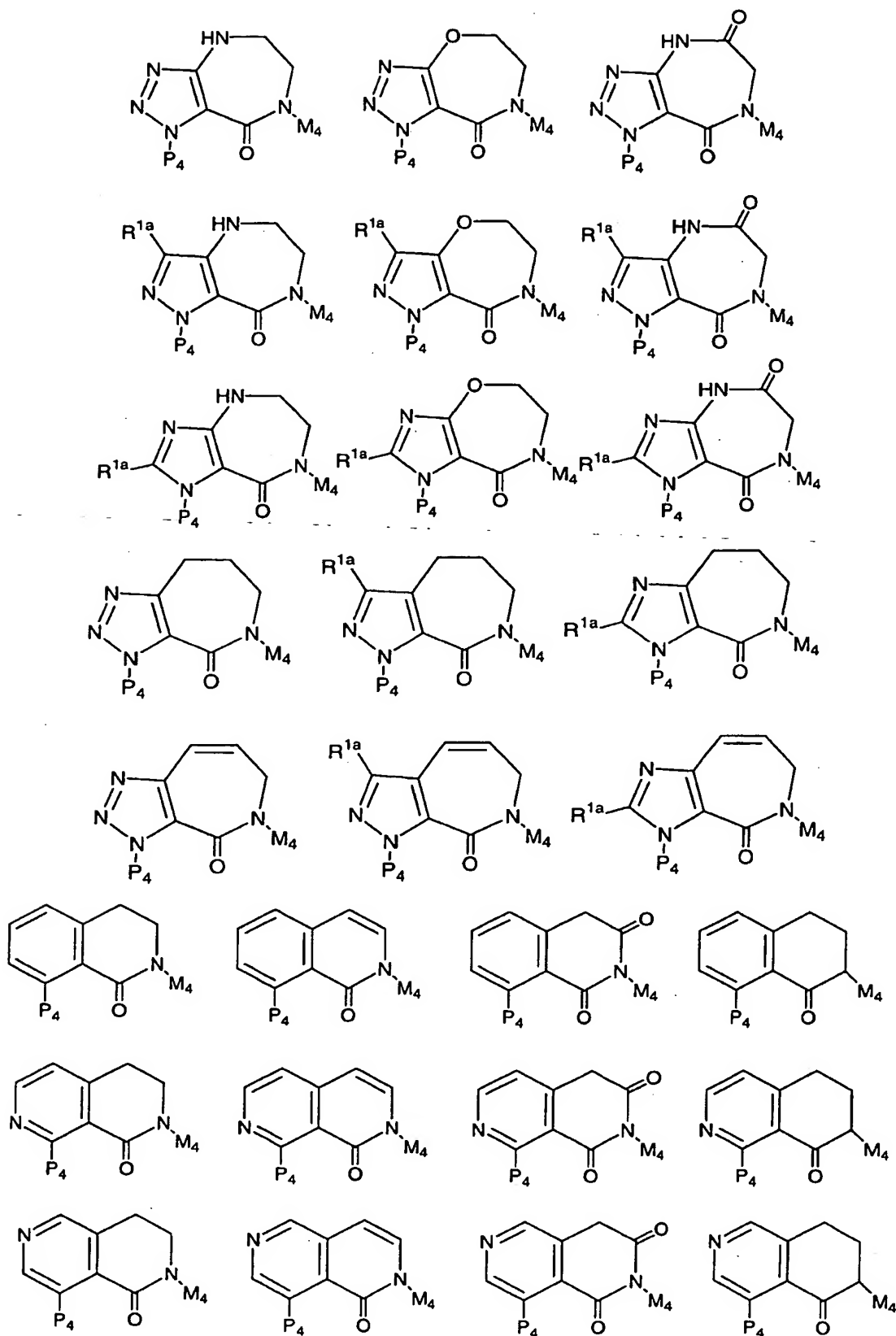
with 0-1 R^{4b}, -CH₂-cyclobutyl substituted with 0-1 R^{4b},
-CH₂-cyclopentyl substituted with 0-1 R^{4b}, benzyl
substituted with 0-2 R^{4b}, 5-6 membered aromatic
heterocycle substituted with 0-2 R^{4b} and consisting of
5 carbon atoms and from 1-4 heteroatoms selected from
the group consisting of N, O, and S(O)_p, and (CH₂)₅₋₆
membered aromatic heterocycle substituted with 0-2 R^{4b}
and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
10 O, and S(O)_p;

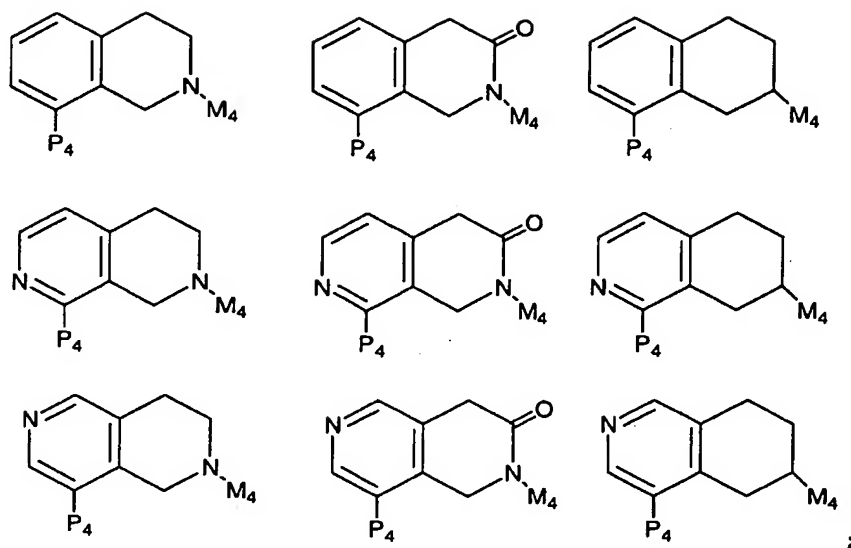
R⁵, at each occurrence, is selected from H, =O, CH₃, CH₂CH₃,
OR³, CH₂OR³, F, Cl, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, C(O)OR^{3c},
NR³C(O)R^{3a}, C(O)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl,
15 NR³SO₂-phenyl, S(O)₂-CH₃, S(O)₂-phenyl, CF₃, phenyl
substituted with 0-2 R⁶, naphthyl substituted with 0-2
R⁶, and benzyl substituted with 0-2 R⁶; and,

R⁶, at each occurrence, is selected from H, OH, OR², F, Cl,
20 CH₃, CH₂CH₃, NR²R^{2a}, CH₂NR²R^{2a}, C(O)R^{2b}, CH₂C(O)R^{2b},
NR²C(O)R^{2b}, and SO₂NR²R^{2a}.

[6] In another preferred embodiment, the present invention
25 provides a novel compound, wherein the compound is selected
from:





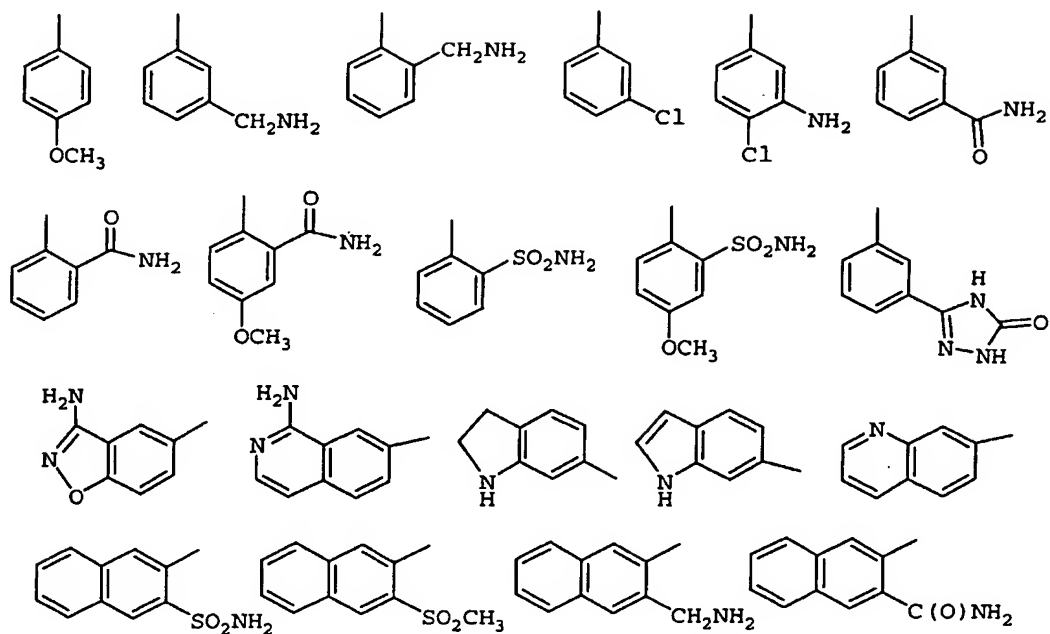


P_4 is $-G_1-G$;

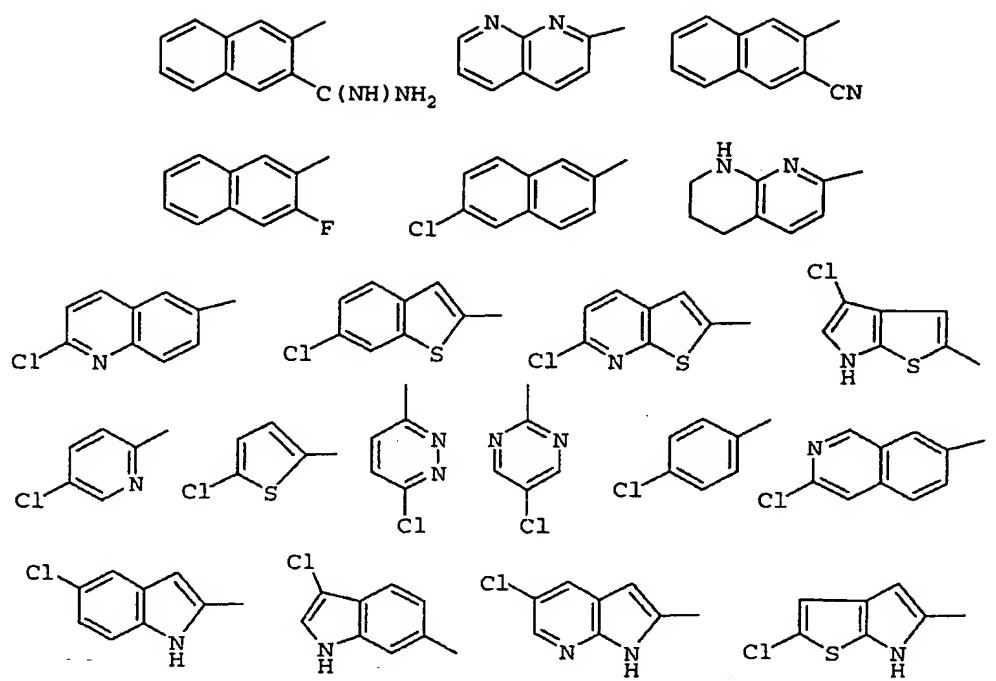
5

M_4 is $-A-B$;

$-G$ is selected from:



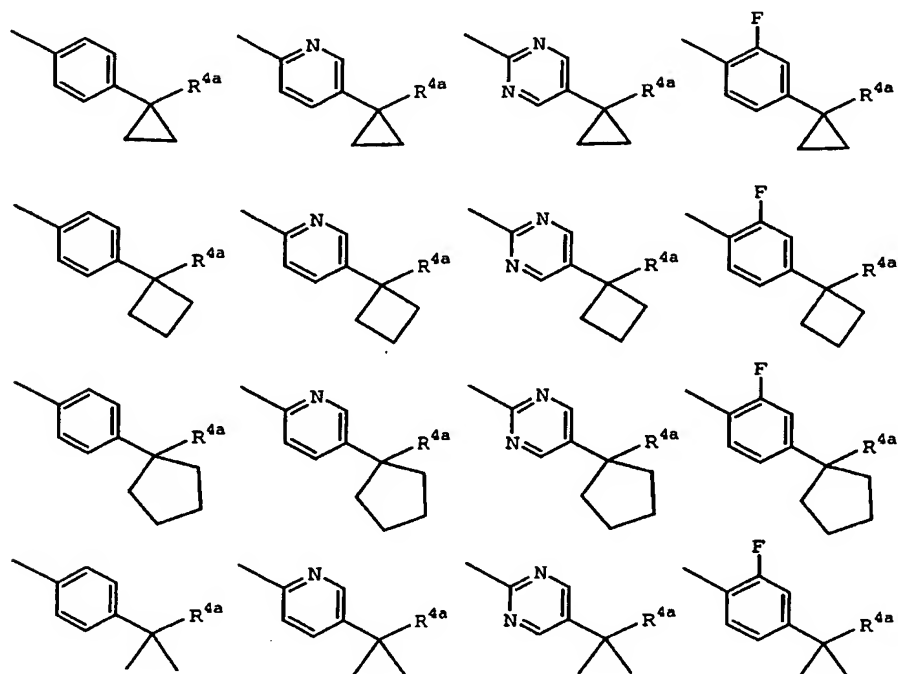
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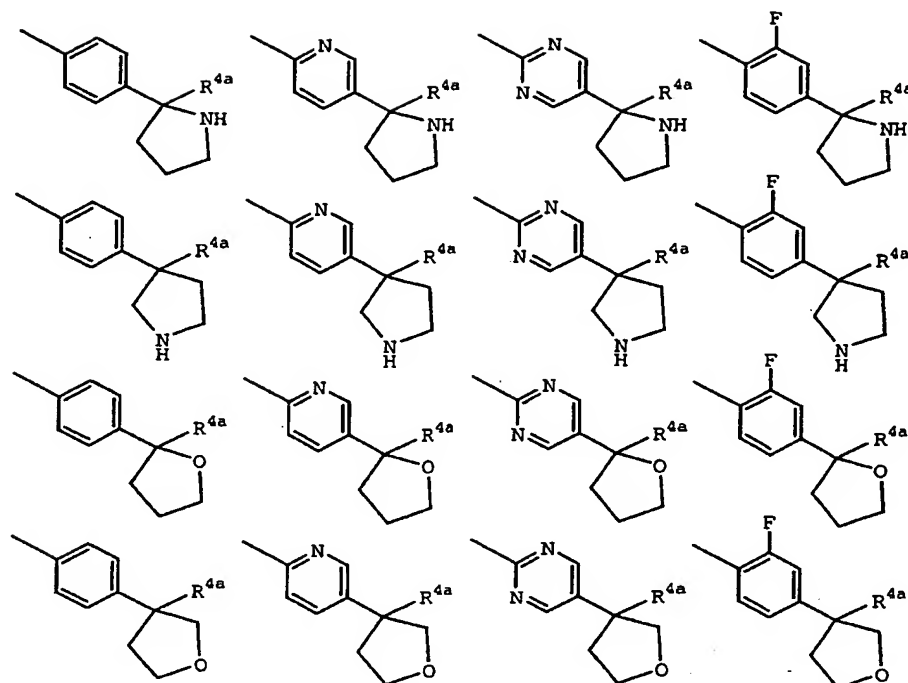


G_1 is absent or is selected from $C(O)NH$, $NHC(O)$, and $NHSO_2$;

5

A-B is selected from:





Z is selected from a bond, CH₂, and CH₂CH₂;

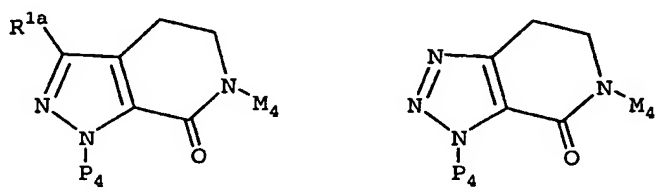
- 5 R^{2d}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl substituted with 0-2 R^{4c}, and a 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;
- 10
- 15 R^{2e}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl, substituted with 0-2 R^{4c}, and 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;
- 20

R^{4a} is selected from $NR^{2d}R^{2d}$, $CH_2NR^{2d}R^{2d}$, $CH_2CH_2NR^{2d}R^{2d}$,
 $N(\rightarrow O)R^{2d}R^{2d}$, $CH_2N(\rightarrow O)R^{2d}R^{2d}$, CH_2OR^{2d} , $C(O)R^{2e}$,
 $C(O)NR^{2d}R^{2d}$, $CH_2C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)R^{2e}$, $CH_2NR^{2d}C(O)R^{2e}$,
5 $NR^{2d}C(O)NR^{2d}R^{2d}$, $CH_2NR^{2d}C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)OR^{2d}$,
 $CH_2NR^{2d}C(O)OR^{2d}$, $NR^{2d}SO_2R^{2d}$, $CH_2NR^{2d}SO_2R^{2d}$, $S(O)_pR^{2d}$,
 $CH_2S(O)_pR^{2d}$, 5-6 membered carbocycle substituted with
0-2 R^{4c} , $-(CH_2)$ -5-6 membered carbocycle substituted
with 0-2 R^{4c} , $-(CH_2)_2$ -5-6 membered carbocycle
10 substituted with 0-2 R^{4c} , 5-6 membered heterocycle
substituted with 0-2 R^{4c} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and $S(O)_p$, $-(CH_2)$ -5-6 membered
heterocycle substituted with 0-2 R^{4c} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the
group consisting of N, O, and $S(O)_p$, and $-(CH_2)_2$ -5-6
membered heterocycle substituted with 0-2 R^{4c} and
consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and $S(O)_p$
20 provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or
 $S(O)H$; and,

R^{4c} is selected from $=O$, OH , OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$,
 $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH=CH_2$,
25 $CH\equiv CH$, CH_2OH , CH_2OCH_3 , $CH_2OCH_2CH_3$, $CH_2OCH_2CH_2CH_3$,
 $CH_2OCH(CH_3)_2$, F , Br , Cl , CF_3 , NR^2R^{2a} , $CH_2NR^2R^{2a}$,
 $C(O)R^{2c}$, $CH_2C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $CH_2NR^2C(O)R^{2b}$,
 $C(O)NR^2R^{2a}$, $CH_2C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $CH_2SO_2NR^2R^{2a}$,
 $NR^2SO_2R^{5a}$, $CH_2NR^2SO_2R^{5a}$, $S(O)_pR^{5a}$, and $CH_2S(O)_pR^{5a}$.

30

[7] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is selected from:

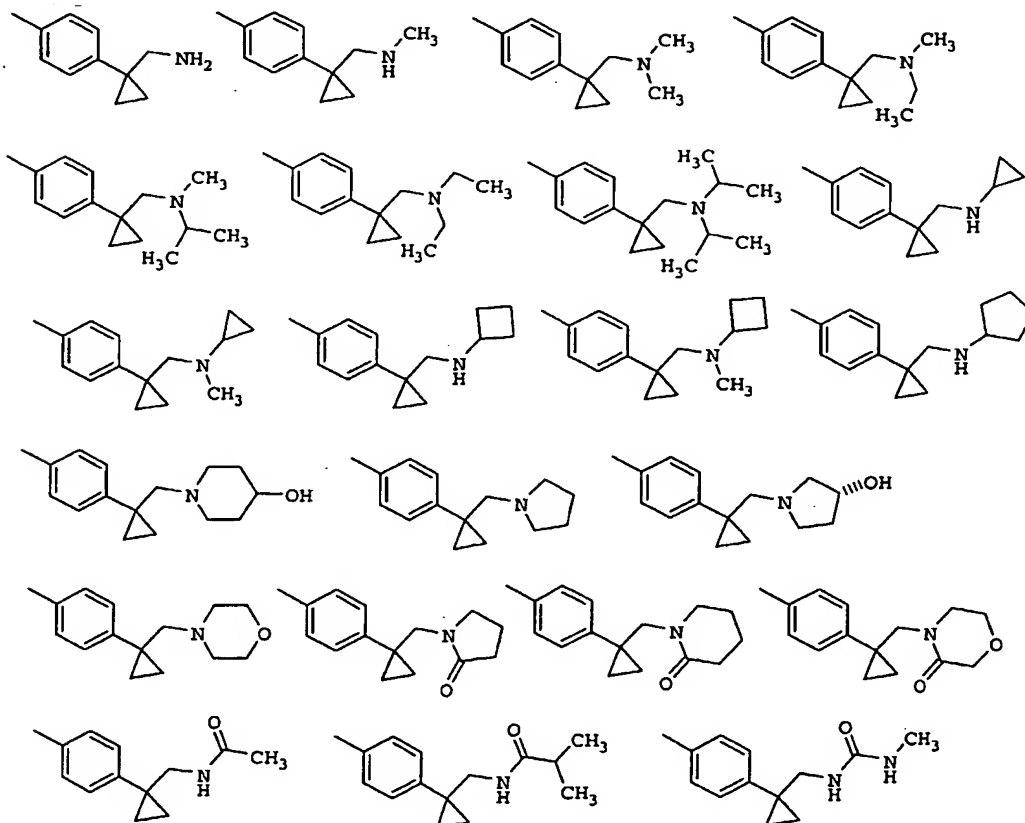


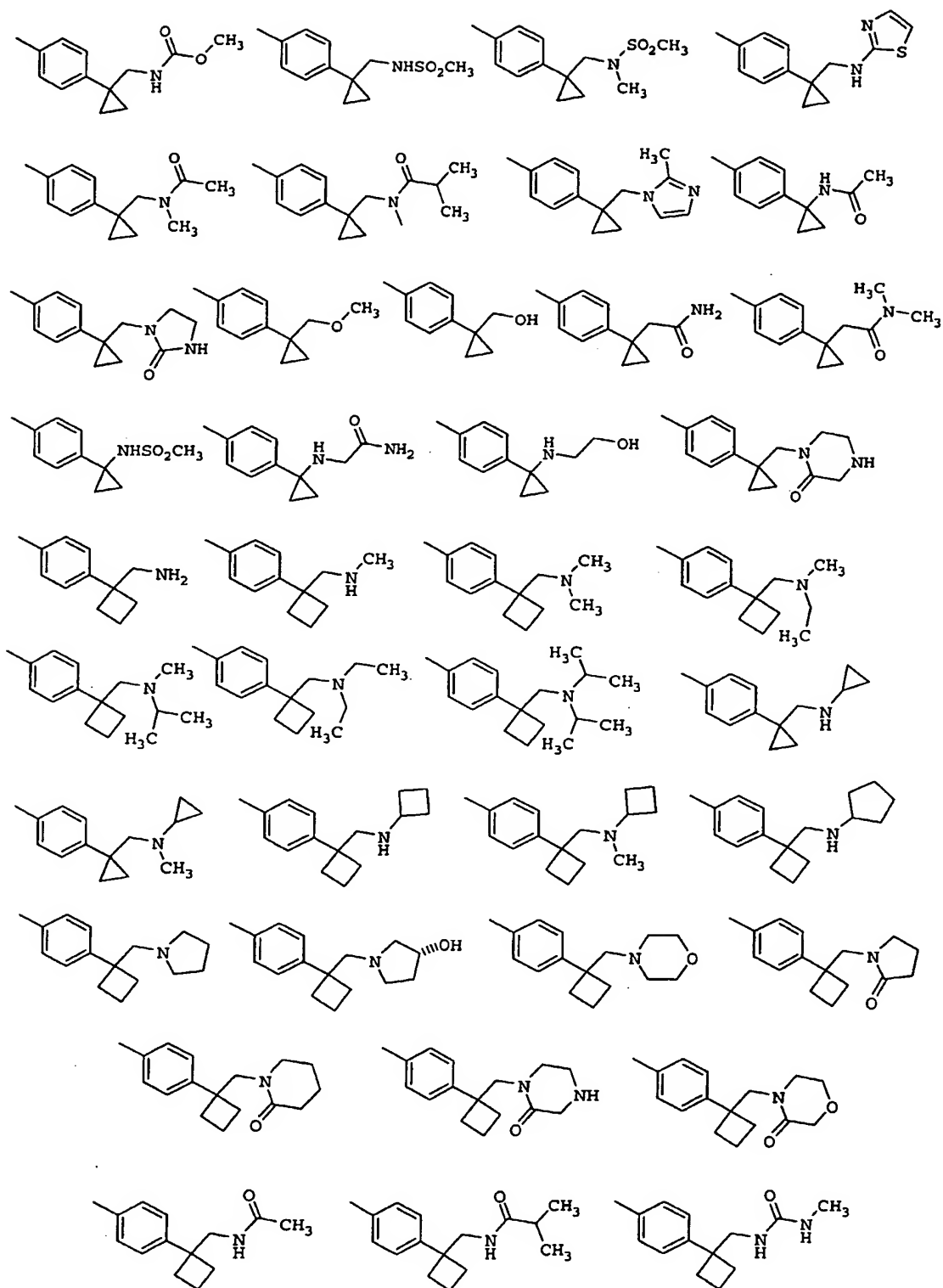
5 P_4 is $-G$;

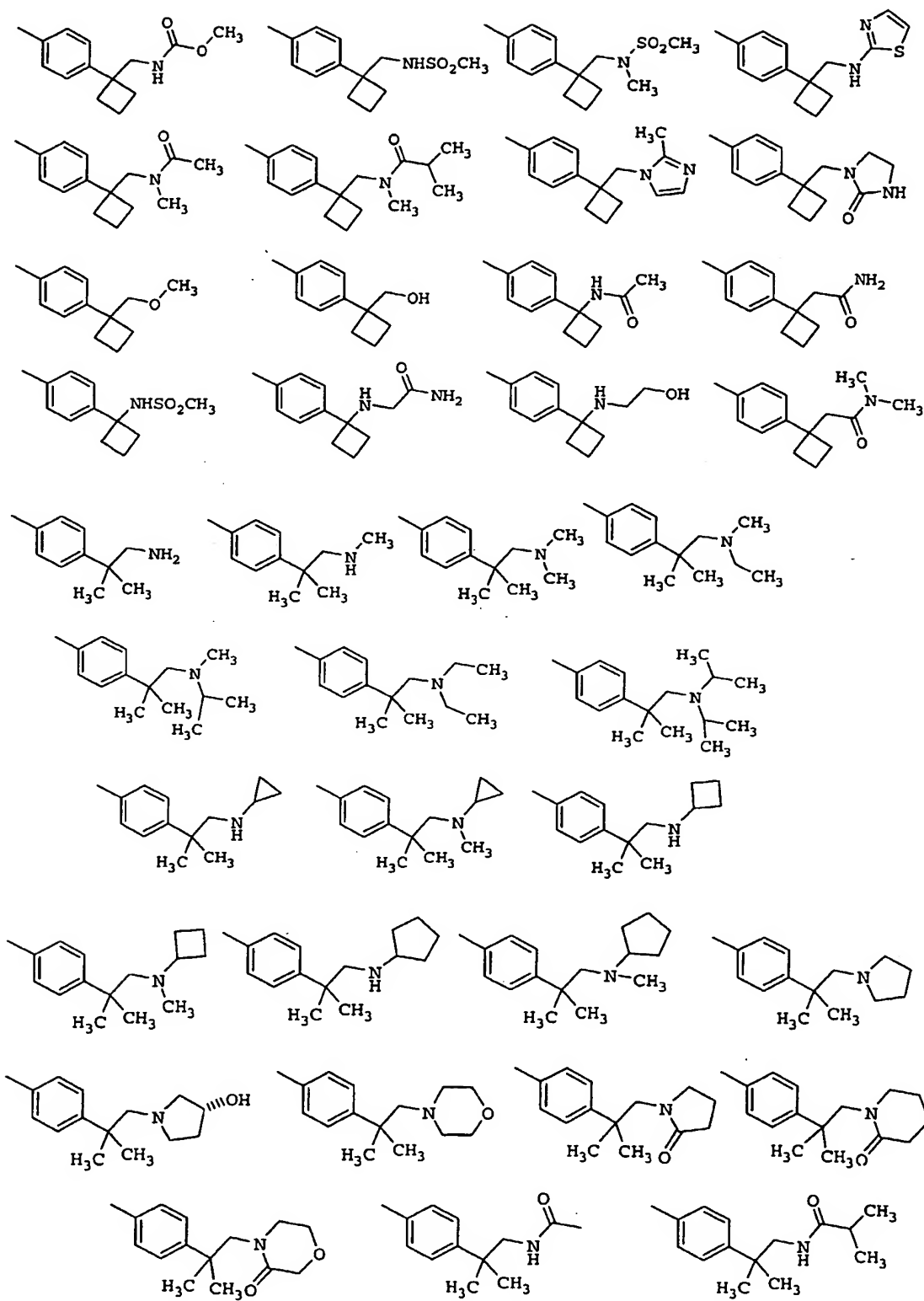
M_4 is $-A-B$;

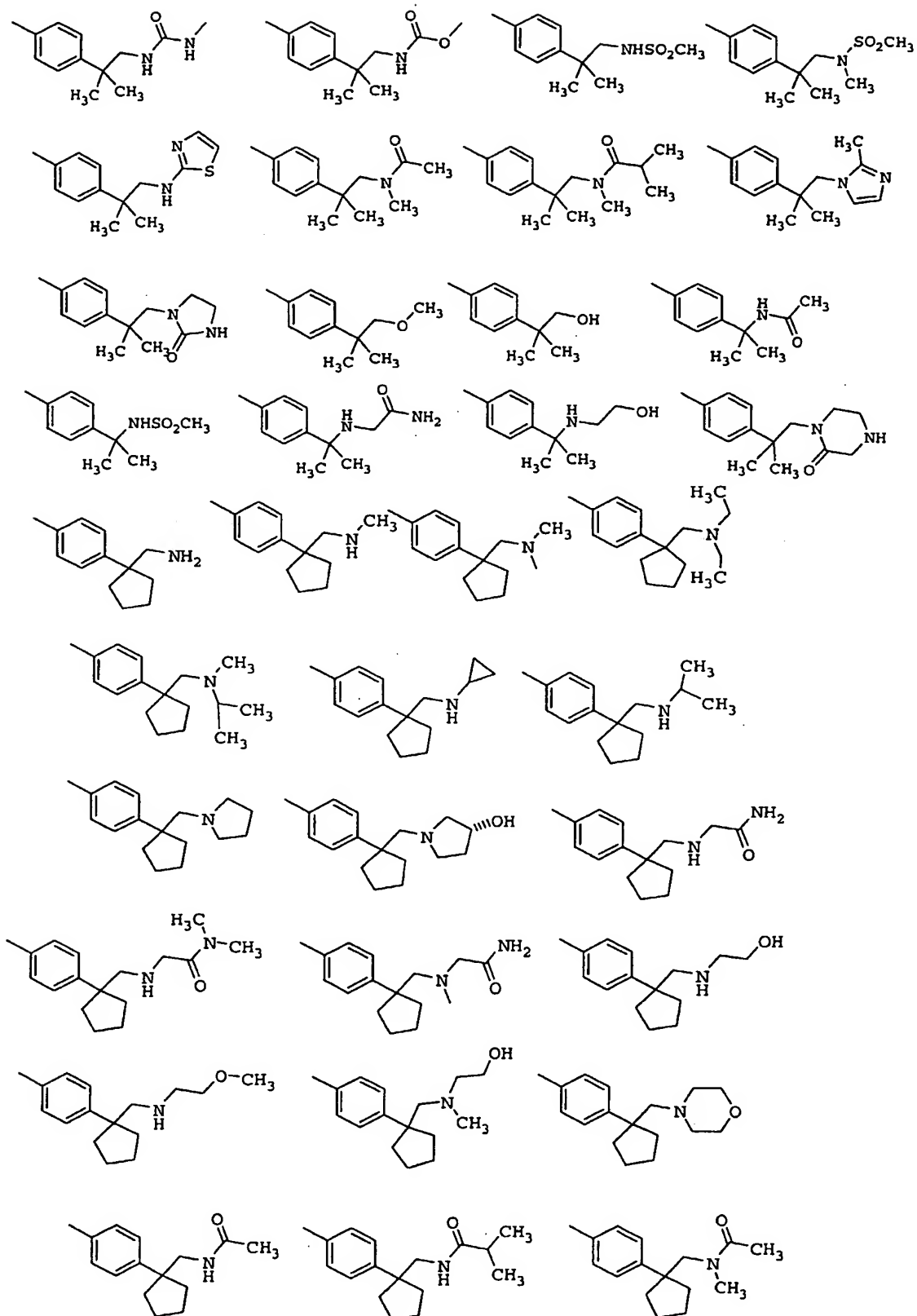
A-B is selected from:

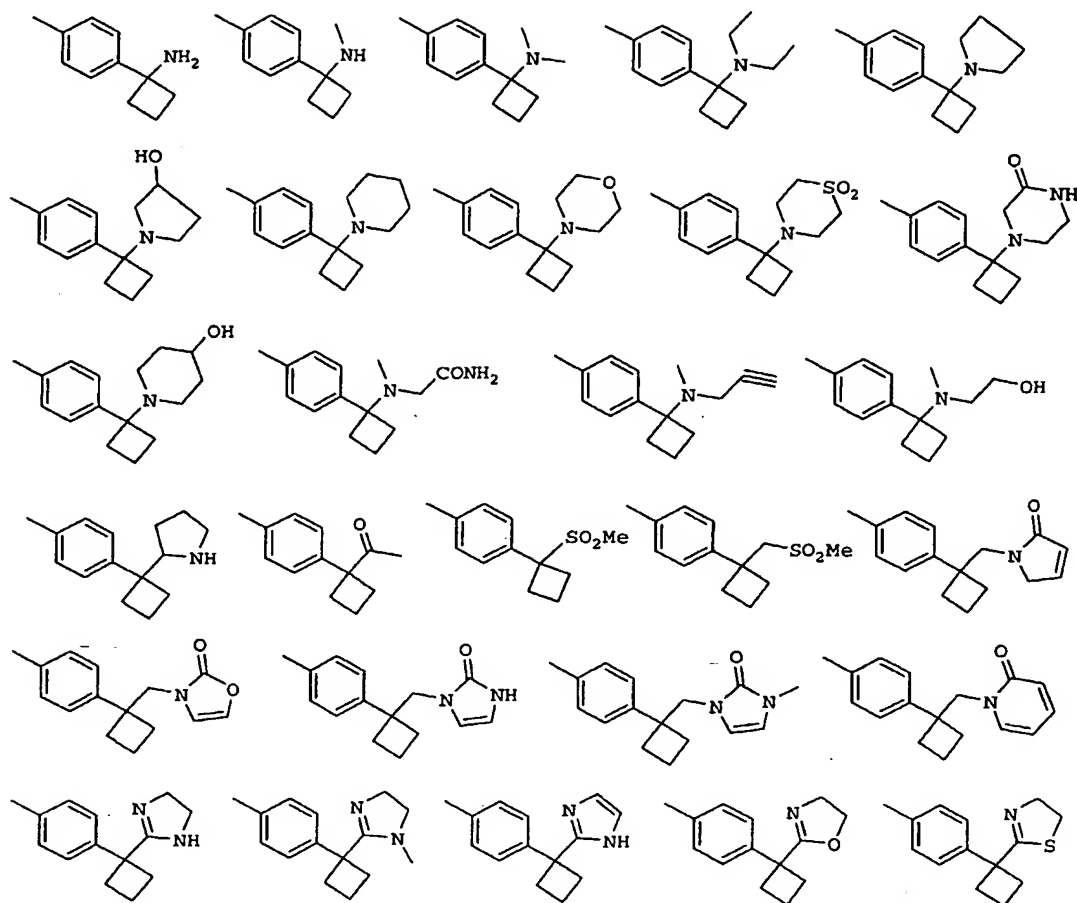
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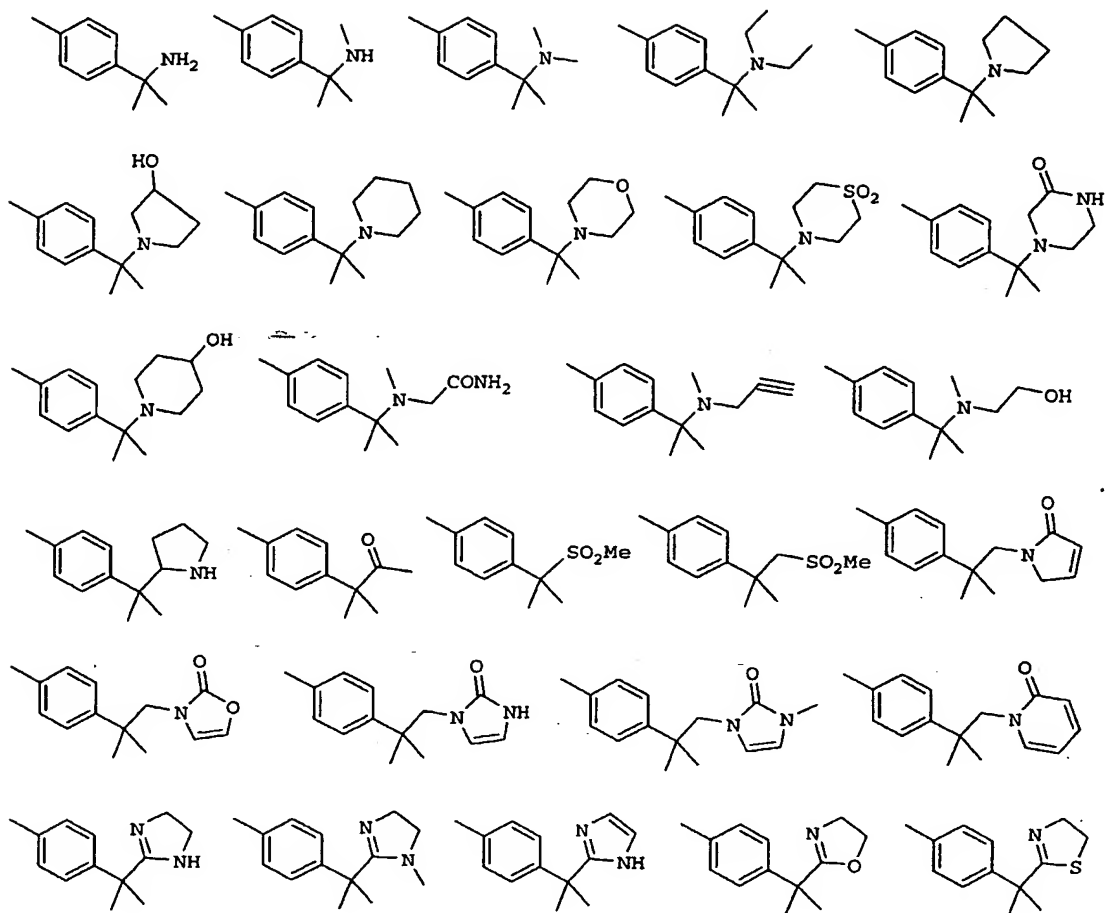


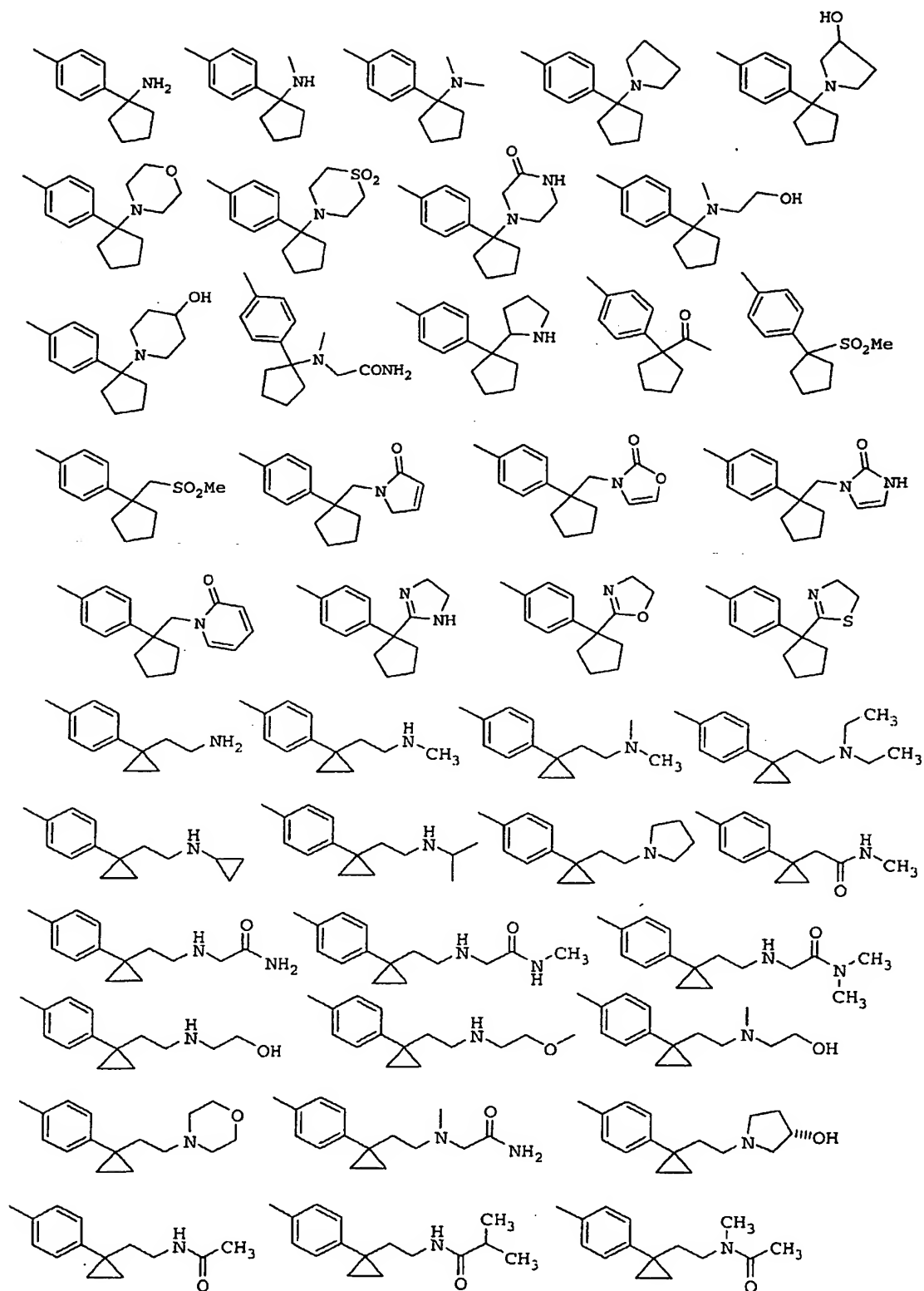


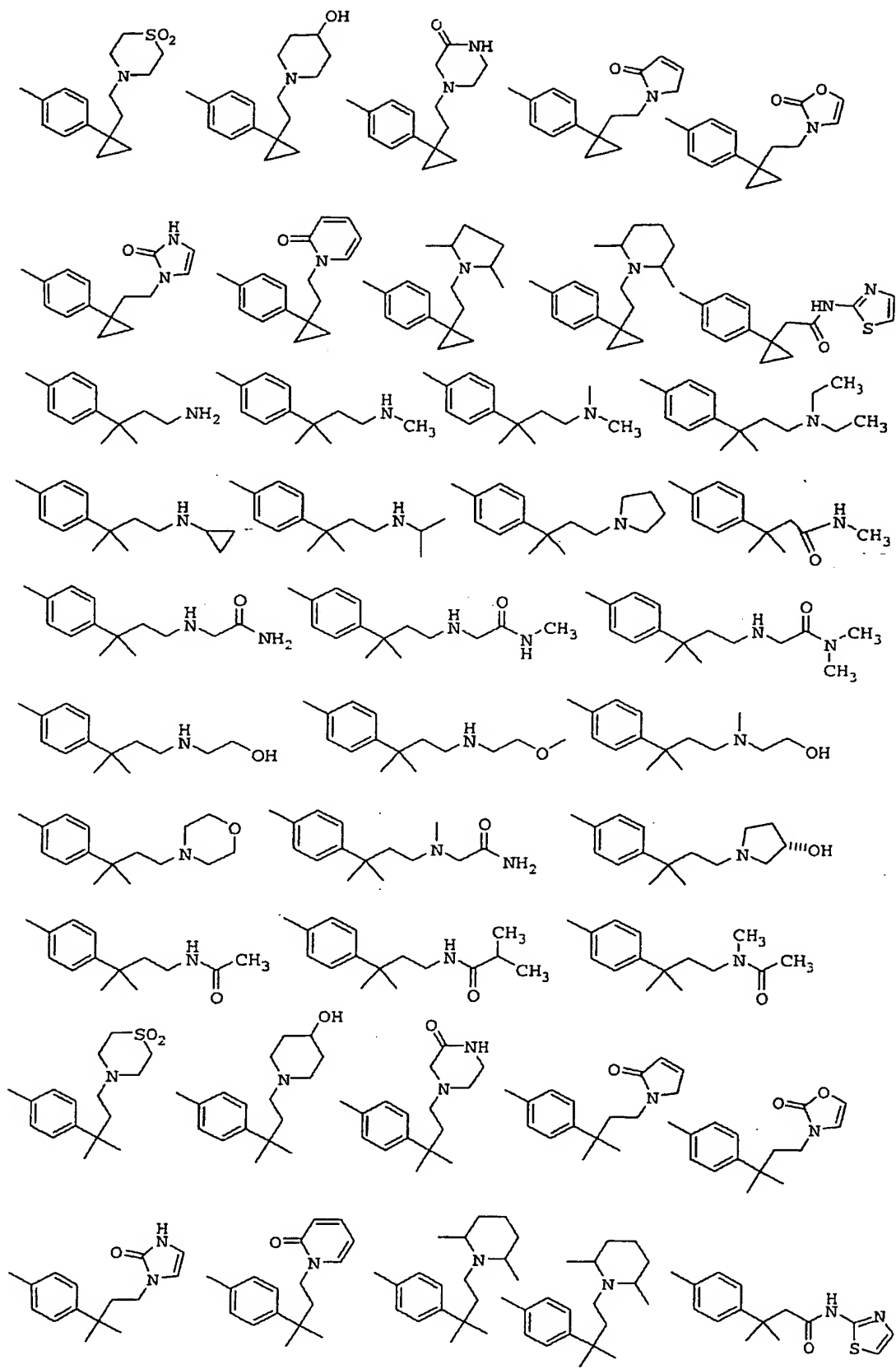


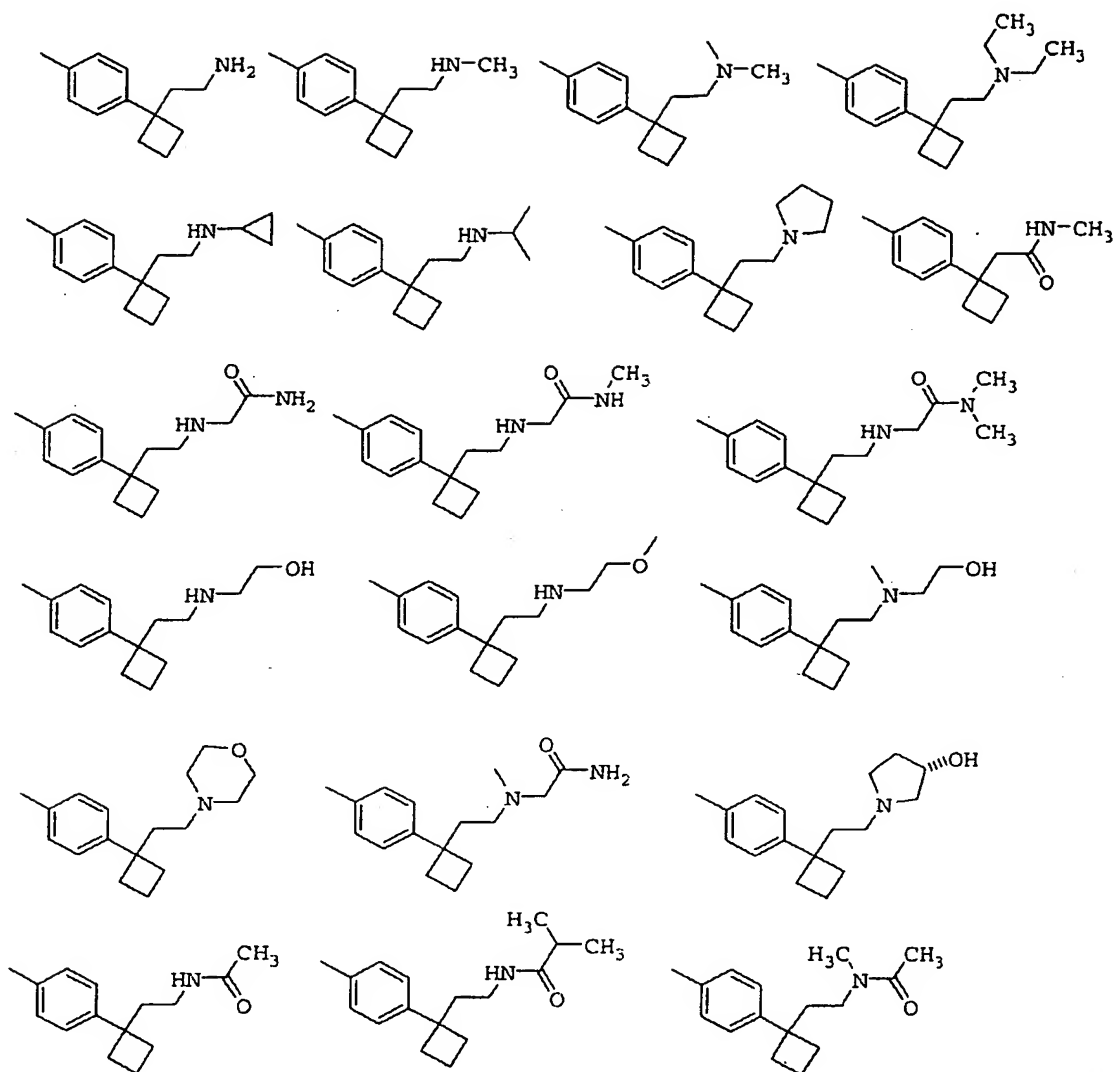


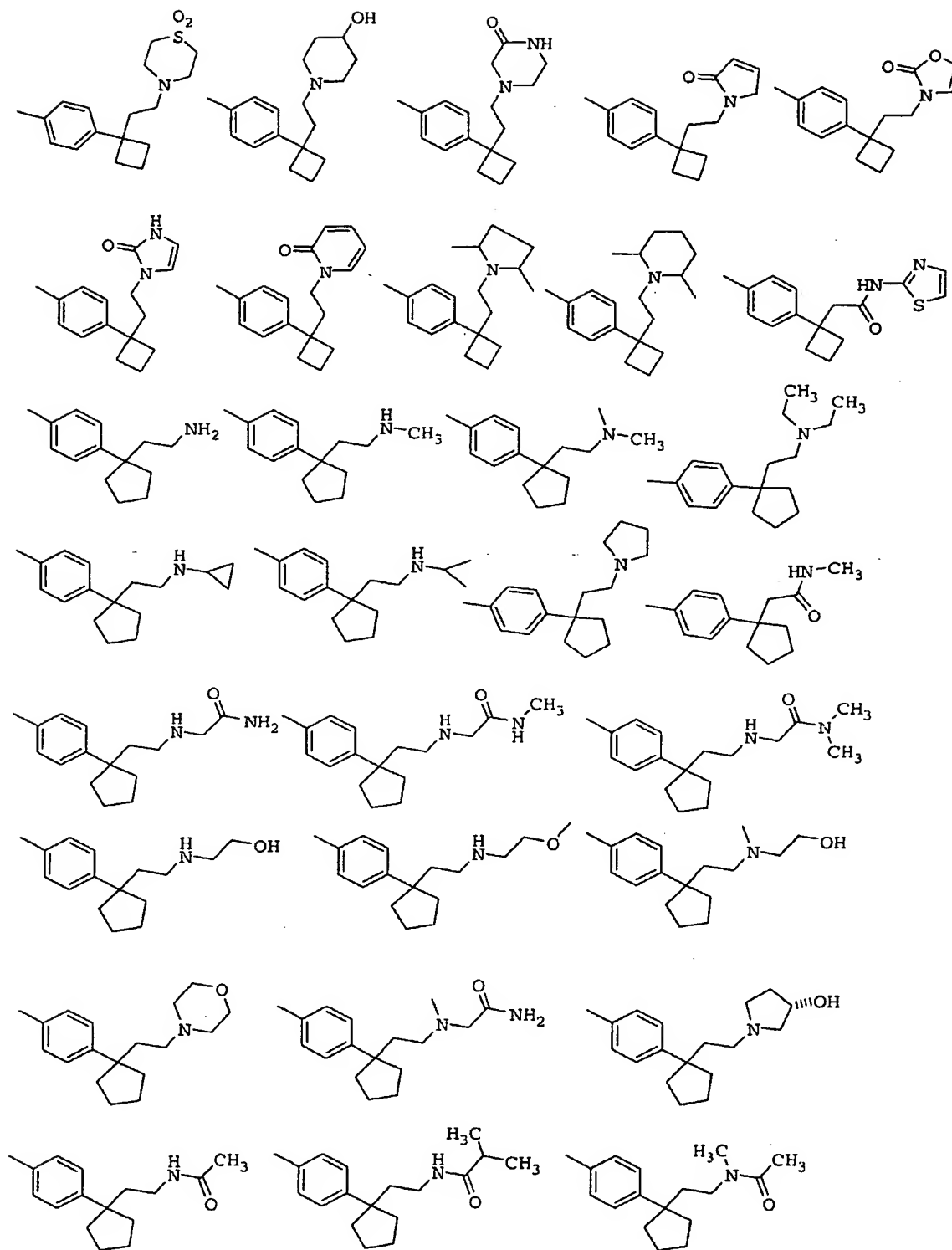


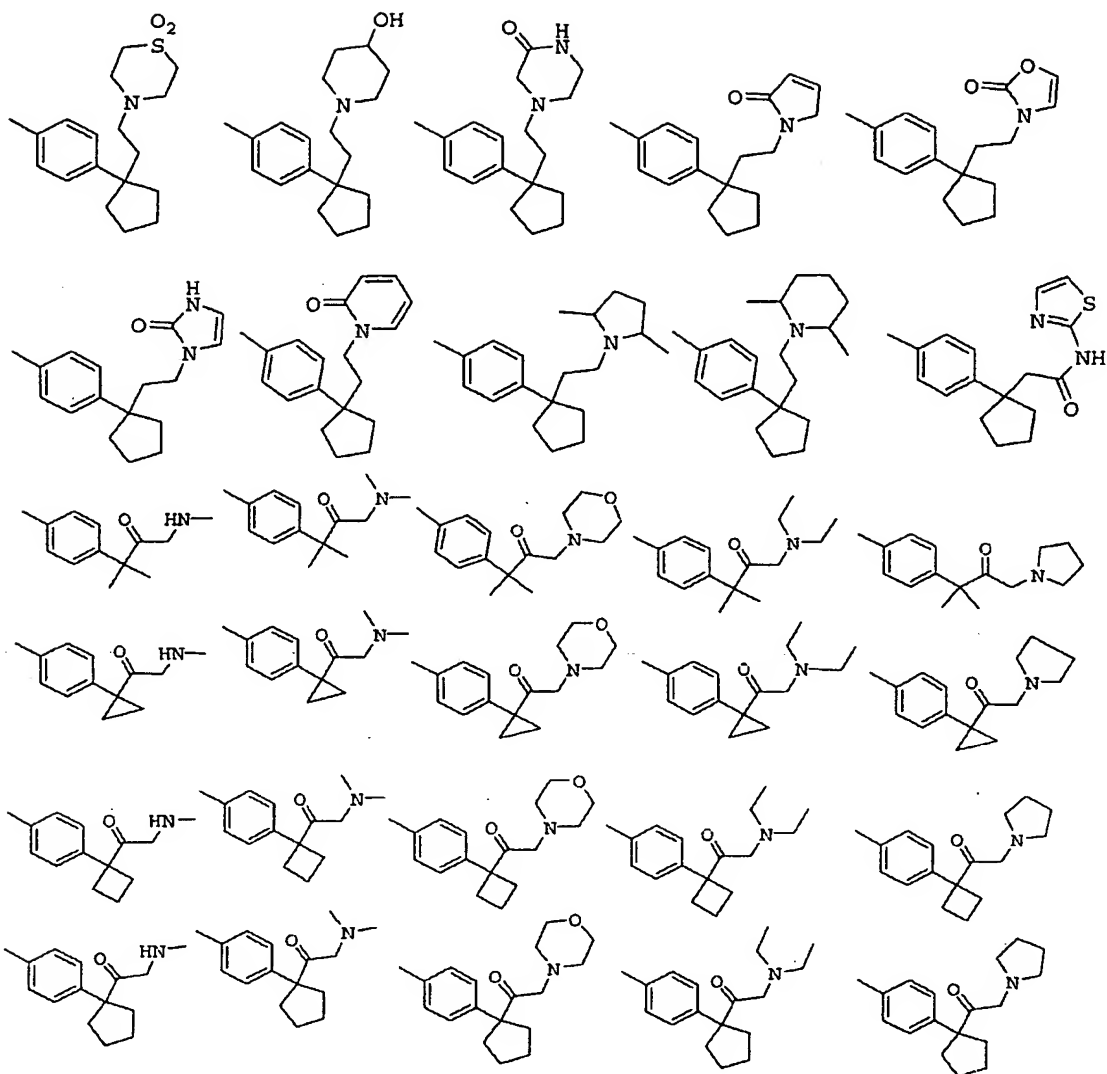












- 5 [8] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is selected from the group:

10 1-(4-methoxyphenyl)-6-(4-(1-
 [(methylamino)methyl]cyclopropyl)phenyl)-3-
 (trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
 c]pyridin-7-one;

- 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-(4-{1-[(cyclopentylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-(4-{1-[(4-hydroxy-1-piperidinyl)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 35 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

- 1-(4-methoxyphenyl)-6-(4-[1-(1-pyrrolidinylcarbonyl)cyclopropyl]phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-N,N-dimethylcyclopropanecarboxamide;
- 10 1-(4-methoxyphenyl)-6-(4-{1-[(4-methyl-1-piperazinyl)carbonyl]cyclopropyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-(4-[1-(4-hydroxypiperidine-1-carbonyl)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropanecarboxamide;
- 25 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropanecarboxylic acid
cyclopentylamide;
- 30 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-N-(1,3,4-thiadiazol-2-yl)cyclopropanecarboxamide;
- 35 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-

yl]phenyl}-N-(1*H*-tetraazol-5-yl)cyclopropanecarboxamide;

5 methyl 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropanecarboxylate;

10 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropanecarbonitrile;

15 6-{4-[1-(aminomethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylacetamide;

20 *N*'-ethyl-*N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylurea;

25 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylethanesulfonamide;

30 1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

35 1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

methyl 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopentanecarboxylate;

5

1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopentyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

10

6-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

15

1-(4-methoxyphenyl)-6-{4-[1-(1-
pyrrolidinylmethyl)cyclopentyl]phenyl}-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

20

6-[4-(1-[(3R)-3-hydroxy-1-
pyrrolidinyl]methyl)cyclopentyl]phenyl]-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

25

1-(4-methoxyphenyl)-6-{4-[1-(4-
morpholinylmethyl)cyclopentyl]phenyl}-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

30

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopentyl)methyl]-N-methylacetamide;

35

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-

yl]phenyl)cyclopentyl)methyl]-N-
methylnmethanesulfonamide;

5 methyl 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl)cyclobutanecarboxylate;

1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclobutyl}phenyl)-3-
10 (trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
15 7H-pyrazolo[3,4-c]pyridin-7-one;

6-(4-{1-[(diethylamino)methyl]cyclobutyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

20 1-(4-methoxyphenyl)-6-{4-[1-(1-
pyrrolidinylmethyl)cyclobutyl]phenyl}-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

25 6-[4-(1-{[(3R)-3-hydroxy-1-
pyrrolidinyl]methyl}cyclobutyl)phenyl]-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

30 1-(4-methoxyphenyl)-6-(4-[1-(4-
morpholinylmethyl)cyclobutyl]phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

35

- N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclobutyl)methyl]-*N*-methylacetamide;
- 5 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclobutyl)methyl]-*N*-methylmethanesulfonamide;
- 10 1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]-phenyl}cyclohexanecarboxylic acid methyl ester;
- 15 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclohexyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;
- 20 6-(4-{1-[(dimethylamino)methyl]cyclohexyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;
- 25 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclohexyl)methyl]-*N*-methylacetamide;
- 30 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclohexyl)methyl]-*N*-methylmethanesulfonamide;
- 35 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7*H*-pyrazolo[3,4-*c*]pyridin-7-one;

- 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-(4-{1-[(cyclopentylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 30 1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 30 N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)methyl]-N-methylacetamide;
- 35 3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

5

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclobutyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7H-pyrazolo[3,4-c]pyridin-7-one;

10

6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

15

6-(4-{1-[(isopropylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

20

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclobutyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

25

6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclobutyl)phenyl]-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

30

6-(4-{1-[(diethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

35

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylacetamide;

5 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

10 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

15 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

20 1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

25 1-(4-methoxyphenyl)-6-(4-{1-(4-morpholinylmethyl)cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

30 6-[4-(1-{[(3*R*)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

35 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-

4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

5 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

10 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

15 1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

20 6-[4-(1-[(3R)-3-hydroxy-1-pyrrolidinylmethyl]cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

25 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

30 1-(3-chlorophenyl)-6-{4-[1-(isopropylamino)methyl]cyclopropyl}phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

35 1-(3-chlorophenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

5

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

10 1-(3-chlorophenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

15 1-(3-chlorophenyl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

20 1-(3-chlorophenyl)-6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

25 1-(3-chlorophenyl)-6-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

30 1-(3-chlorophenyl)-6-(4-{1-[(cyclopropylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

35 1-(3-chlorophenyl)-6-(4-{1-[(cyclobutylamino)methyl]cyclopropyl}phenyl)-7-oxo-

4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

5 1-(3-chlorophenyl)-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

10 1-(3-chlorophenyl)-6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

15 1-(3-chlorophenyl)-6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

20 1-(3-chlorophenyl)-6-{4-[1-(methoxymethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-6-{4-[1-(methoxymethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

25 1-(3-chlorophenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

30 1-(3-chlorophenyl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

35 1-(3-chlorophenyl)-6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-

7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-
5 pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-
tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

N-[(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
10 yl]phenyl}cyclopropyl)methyl]-N-methylacetamide;

6-(4-{1-[(cyclopropylamino)methyl]cyclopropyl}phenyl)-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;

15 6-(4-{1-[(cyclobutylamino)methyl]cyclopropyl}phenyl)-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;

20 6-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-
1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;

6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}
25 cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-
tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

6-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
30 pyrazolo[3,4-c]pyridine-3-carbonitrile;

5-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-3-(4-
methoxyphenyl)-3,5,6,7-tetrahydro-4H-
[1,2,3]triazolo[4,5-c]pyridin-4-one;

35

5-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-3-(4-methoxyphenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

5 3-(4-methoxyphenyl)-5-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

3-(4-methoxyphenyl)-5-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

3-(4-methoxyphenyl)-5-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

5-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-3-(4-methoxyphenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

20 3-(3-chlorophenyl)-5-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

25 3-(3-chlorophenyl)-5-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

30 3-(3-chlorophenyl)-5-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

3-(3-chlorophenyl)-5-(4-{1-[(3-hydroxy-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

6-[4-(1-[(2-hydroxyethyl)(methylamino)methyl]cyclopropyl)phenyl]-
1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
5 pyrazolo[3,4-c]pyridine-3-carboxamide;

6-(4-[1-(dimethylamino)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxamide;

10

6-(4-{1-[(2-hydroxyethyl)(methylamino)cyclopropyl]phenyl}-
1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxamide;

15 2-(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;

6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-
20 1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxamide;

2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
25 yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;

2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)acetamide;

30

2-(1-{4-[1-(3-chlorophenyl)-3-cyano-7-oxo-1,4,5,7-
tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)acetamide;

- 1-(3-chlorophenyl)-6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 2-(1-{4-[1-(3-chlorophenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 10 2-(1-{4-[3-(3-chlorophenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 15 2-(1-{4-[3-(4-methoxyphenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 20 2-(1-{4-[3-(4-methoxyphenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)acetamide;
- 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-imidazolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-piperazinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(3-oxo-4-morpholinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 35 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-piperidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 2-[(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)oxy]acetamide;
- 5 1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl carbamate;
- 10 2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)acetamide;
- 15 2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(methylamino)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 6-{4-[1-(dimethylamino)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1,3-thiazol-2-ylamino)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 N-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)urea;
- 35 N-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N'-methylurea;

N-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)-2-methylpropanamide;

5 6-(4-{1-[(4-hydroxy-1-piperidinyl)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

10 1-(4-methoxyphenyl)-6-(4-{1-[(2-methyl-5,6-dihydro-1(4*H*)-pyrimidinyl)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

15 1-(4-methoxyphenyl)-6-(4-{1-[(2-methyl-4,5-dihydro-1*H*-imidazol-1-yl)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

20 6-{4-[1-(4,5-dihydro-1,3-oxazol-2-ylmethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

25 6-{4-[1-(4,5-dihydro-1*H*-imidazol-2-ylmethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

30 1-(4-methoxyphenyl)-6-(4-{1-[(1-methyl-4,5-dihydro-1*H*-imidazol-2-yl)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

35 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(1,3-thiazol-2-ylamino)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

- 1- (4-methoxyphenyl) -6- (4- {1- [(2-methyl-1*H*-imidazol-1-yl)methyl]cyclopropyl}phenyl) -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;
- 5 1- (4-methoxyphenyl) -6- {4- [1-methyl-1- (2-oxo-1-pyrrolidinyl)ethyl]phenyl} -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 10 1- (4-methoxyphenyl) -6- {4- [1-methyl-1- (2-oxo-1-piperidinyl)ethyl]phenyl} -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 15 6- {4- [1,1-dimethyl-2- (2-oxo-1-piperidinyl)ethyl]phenyl} -1- (4-methoxyphenyl) -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 20 6- {4- [1,1-dimethyl-2- (2-oxo-1-pyrrolidinyl)ethyl]phenyl} -1- (4-methoxyphenyl) -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 25 6- {4- [1,1-dimethyl-2- (3-oxo-4-morpholinyl)ethyl]phenyl} -1- (4-methoxyphenyl) -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 30 6- {4- [1,1-dimethyl-2- (2-oxotetrahydro-1 (2*H*) -pyrimidinyl)ethyl]phenyl} -1- (4-methoxyphenyl) -7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

- 6-{4-[1,1-dimethyl-2-(2-oxodihydro-2*H*-1,3-oxazin-3(4*H*)-
yl)ethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-
tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 5 1-{4-[3-(aminocarbonyl)-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-1-
methylethyl methylcarbamate;
- 10 1-{4-[3-(aminocarbonyl)-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-1-
methylethyl 3-pyrrolidinylcarbamate;
- 15 6-{4-[1-ethyl-1-(1-pyrrolidinylmethyl)propyl]phenyl}-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-
pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 20 6-(4-{1-[(dimethylamino)methyl]-1-ethylpropyl}phenyl)-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-
pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 1-[3-(aminomethyl)phenyl]-6-{4-[1,1-dimethyl-2-(1-
pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-
1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 25 1-[3-(aminomethyl)phenyl]-6-{4-[2-(dimethylamino)-1,1-
dimethylethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-
pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 30 1-[3-(aminocarbonyl)phenyl]-6-{4-[2-(dimethylamino)-1,1-
dimethylethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-
pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 35 1-[3-(aminocarbonyl)phenyl]-6-{4-[1,1-dimethyl-2-(1-
pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-
1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

- 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 10 1-(1-amino-7-isoquinolinyl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 1-(1-amino-7-isoquinolinyl)-6-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 1-(1-amino-7-isoquinolinyl)-6-{4-[1-[(dimethylamino)methyl]cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 25 1-(1-amino-7-isoquinolinyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 30 1-(3-amino-1,2-benzisoxazol-5-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

- 1-(3-amino-1,2-benzisoxazol-5-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-[3-(aminomethyl)phenyl]-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 10 6-[4-(1-{[acetyl(methyl)amino]methyl}cyclopropyl)phenyl]-1-[3-(aminomethyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 1-[3-(aminocarbonyl)phenyl]-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 3-[3-cyano-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridin-1-yl]benzamide;
- 25 1-(2,3-dihydro-1H-indol-6-yl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 30 1-(2,3-dihydro-1H-indol-6-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 1-(2,3-dihydro-1H-indol-6-yl)-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

- 6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 10 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopentyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 6-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(2-oxo-pyrrolidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 1-(4-methoxyphenyl)-6-{4-[1-(2-oxo-piperidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 N-(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-cyclopropyl)-N-methyl-acetamide;

N-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-methanesulfonamide;

5 *N*-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-2-methylaminoacetamide;

10 2-dimethylamino-*N*-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)-*N*-methylacetamide;

15 *N*-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-2-morpholin-4-yl-acetamide;

20 6-{4-[1-(1-hydroxyethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

25 6-[4-(1-acetylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

30 6-{4-[1-(1-hydroxy-1-methyl-ethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

6-[4-(1-methoxymethylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

- 6-{4-[1-(4,5-dihydro-oxazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropanecarboxylic acid 2-amino-ethyl ester ;
- 10 6-{4-[1-(4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 6-{4-[1-(4,5-dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(1-methyl-4,5-dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-{4-[1-(1-methanesulfonyl-4,5-dihydro-1H-imidazol-2-yl)-cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-{4-[1-(1H-imidazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 1-(4-methoxyphenyl)-6-{4-[1-(1-methyl-1H-imidazol-2-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 2-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-cyclopropyl)-methyl-amino]-acetamide;

6-(4-{1-[(2-hydroxyethyl)-methylamino]cyclopropyl}phenyl)-
1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-
tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

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1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopropanecarboxylic acid methoxy-methyl-amide;

10 6-[4-(1-hydroxymethylcyclopropyl)phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amide;

15 6-[4-(1-acetyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-
oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide ;

20 6-[4-(1-aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide;

25 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)-
phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amid;

25

6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

30 6-[4-(1-methylaminomethylcyclopentyl)phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4c]pyridine-3-carboxylic acid amide;

35 6-[4-(1-dimethylaminomethylcyclopentyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

6-[4-(1-dimethylaminomethylcyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

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6-[4-(1-[(2-hydroxyethyl)methylaminomethyl]cyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

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6-[4-(1-hydroxymethylcyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15 6-(4-{1-[(2-hydroxyethyl)methylamino]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

20 1-(4-methoxyphenyl)-6-{4-[1-(methyl-prop-2-ynylamino)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

25 3-(1-hydroxyethyl)-1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

3-acetyl-1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

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1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid methylamide;

- 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid dimethylamide;
- 5 6-[4-(1-aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carbonitrile;
- 10 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carbonitrile;
- 15 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 20 2-[(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-
yl}phenyl)cyclopropyl)-methylamino]acetamide;
- 25 6-(4-{1-[(2-hydroxyethyl)methylamino]cyclopropyl}phenyl)-1-
(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 1-(4-methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-
cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 35 1-(4-methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-
cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 1-(4-methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 5 1-(4-methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid methylamide;
- 15 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid dimethylamide;
- 20 6-[4-[1-(1,1-dioxo-1 λ ⁶-thiomorpholin-4-yl)cyclopropyl]phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-aminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 6-[4-(1-dimethylaminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(3-chloro-phenyl)-6-[4-[1,1-dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

- 6-{4-[1,1-dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-(4-methoxy-phenyl)-6-[4-(1-methyl-1-pyrrolidin-1-ylethyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 10 6-[4-(1-dimethylamino-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 6-{4-[1-(4,4-dimethyl-4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-[4-(1-methanesulfonyl-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-hydroxy-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 35 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;

- 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 5 1-(4-methoxy-phenyl)-6-{4-[1-(2-oxo-2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-acetyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;

- 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 5 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 10 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 20 6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 1-(4-methoxy-phenyl)-6-[4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 35 1-(4-methoxy-phenyl)-7-oxo-6-(4-[1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl]-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-[4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

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6-{4-[1-(2-diethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

10 1-(4-methoxy-phenyl)-7-oxo-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15 6-(4-{1-[2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

20 6-(4-{1-[2-(3-hydroxy-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

25 6-(4-{1-[2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

30 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-piperidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

35 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

1-(4-methoxy-phenyl)-6-(4-{1-[2-(methyl-thiazol-2-yl-amino)-ethyl]-cyclopropyl}-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

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6-[4-(1-{2-[(2-hydroxy-ethyl)-methyl-amino]-ethyl}-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

10

1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-methyl-imidazol-1-yl)-ethyl]-cyclopropyl}-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15

6-(4-{1-[2-(2,6-dimethyl-piperidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

20

2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;

25

2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;

30

2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;

35

2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;

- 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 3-methanesulfonyl-6-{4-[1-(2-methoxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-diethylamino-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-(4-{1-[2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

6-(4-{1-[2-(3-hydroxy-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

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3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-piperidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

10 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

15 6-[4-(1-{2-[(2-hydroxy-ethyl)-methyl-amino]-ethyl}-cyclopropyl)-phenyl]-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

20 2-([2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethyl]-methyl-amino)-acetamide;

25 2-[2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethylamino]-acetamide;

30 6-(4-{1-[2-(2-hydroxy-ethylamino)-ethyl]-cyclopropyl}-phenyl)-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-methyl-imidazol-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(thiazol-2-ylamino)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 15 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;
- 20 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 35 1-(4-methoxy-phenyl)-7-oxo-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 1-(4-methoxy-phenyl)-6-(4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 6-(4-(1-[2-(1,1-dioxo-1,6-thiomorpholin-4-yl)-ethyl]-cyclopropyl)-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-(4-(1-[2-(2-hydroxy-ethylamino)-ethyl]-cyclopropyl)-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 15 2-[2-(1-(4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropyl)-ethylamino]-acetamide;
- 20 2-([2-(1-(4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropyl)-ethyl)-methyl-amino]-acetamide;
- 25 6-[4-(1-(2-[(2-hydroxy-ethyl)-methyl-amino]-ethyl)-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 N-[2-(1-(4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropyl)-ethyl]-N-methyl-methanesulfonamide;
- N-[2-(1-(4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropyl)-ethyl]-N-methyl-acetamide;

- 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-methyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 5-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 25 5-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 30 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 10 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-3-methyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 5-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;

- 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-N-methyl-acetamide;
- 5 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-N,N-dimethyl-acetamide;
- 10 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-acetamide;
- 15 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-
phenyl}-cyclopentyl)-acetamide;
- 20 6-[4-(1-carbamoylmethyl-cyclopentyl)-phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-
cyclopentyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-dimethylcarbamoylmethyl-cyclopentyl)-phenyl]-1-(4-
methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopentyl)-N,N-dimethyl-acetamide;
- 35 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopentyl)-N-methyl-acetamide;

- 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-N-methyl-acetamide;
- 5 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-N,N-dimethyl-acetamide;
- 10 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-acetamide;
- 15 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;
- 20 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 25 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N,N-dimethyl-acetamide;
- 30 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 35 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;

- 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;
- 5 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 10 2-(1-{4-[1-(4-methoxy-phenyl)-3-methyl-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 15 2-(1-{4-[1-(4-methoxy-phenyl)-3-methyl-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;
- 20 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;
- 25 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N,N-dimethyl-acetamide;
- 30 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-cyclobutyl)-N,N-dimethyl-acetamide;
- 35 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;

- 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-cyclobutyl)-acetamide;
- 5 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 10 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 15 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 20 5-chloro-thiophene-2-carboxylic acid [2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 25 5-chloro-thiophene-2-carboxylic acid [2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 30 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 35 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;

- 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid {2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid (2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 5 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 10 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 15 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 20 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 25 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 30 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

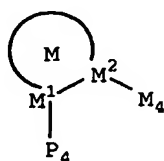
- 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetic acid;
- 2- (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 25 2- (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;
- 30 2- (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 35 1- (4-methoxy-phenyl)-6- {4-[1-(2-oxo-2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 6-(4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl)-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-(4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl)-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one
- 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-acetyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 30 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 35 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 1-(4-Methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

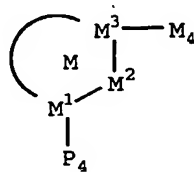
- 6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 5 6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 10 6-[4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 1-(4-methoxy-phenyl)-6-[4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide; and,
- 20 1-(4-methoxy-phenyl)-6-[4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

or a pharmaceutically acceptable salt form thereof.

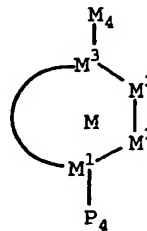
- 25 [9] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is of Formula IIIa, IIIb, or IIIc:



IIIa



IIIb



IIIc

- 30 or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

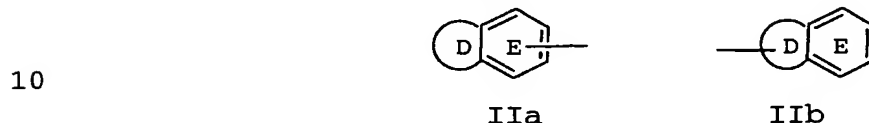
ring M, including M₁, M₂, and, if present, M₃, is phenyl or a 3-10 membered carbocyclic or 4-10 membered

heterocyclic ring consisting of: carbon atoms and 1-4 heteroatoms selected from O, S(O)_p, N, and NZ²;

ring M is substituted with 0-3 R^{1a} and 0-2 carbonyl groups,
5 and there are 0-3 ring double bonds;

one of P₄ and M₄ is -Z-A-B and the other -G₁-G;

G is a group of formula IIa or IIb:



ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon
15 atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;
20

E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

alternatively, ring D is absent, and ring E is selected
25 from phenyl, pyridyl, pyrimidyl, and thienyl, and ring E is substituted with 1-3 R;

alternatively, ring D is absent, ring E is selected from phenyl, pyridyl, and thienyl, and ring E is
30 substituted with 1 R and substituted with a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, wherein the 5-6 membered heterocycle

is substituted with 0-2 carbonyls and 1-3 R and there are 0-3 ring double bonds;

R is selected from H, C₁₋₄ alkyl, F, Cl, OH, OCH₃, OCH₂CH₃,
 5 OCH(CH₃)₂, CN, C(=NH)NH₂, C(=NH)NHOH, C(=NH)NHOCH₃,
 NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂,
 CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tNR⁷R⁸,
 C(O)NR⁷R⁸, CH₂C(O)NR⁷R⁸, S(O)_pNR⁷R⁸, CH₂S(O)_pNR⁷R⁸,
 SO₂R³, and OCF₃;

10

alternatively, when 2 R groups are attached to adjacent atoms, they combine to form methylenedioxy or ethylenedioxy;

15 A is selected from:

C₅₋₁₀ carbocycle substituted with 0-2 R⁴, and
 5-10 membered heterocycle substituted with 0-2 R⁴ and
 consisting of: carbon atoms and 1-4 heteroatoms selected
 from the group consisting of N, O, and S(O)_p;

20

X is selected from -(CR²R^{2a})₁₋₄-, -C(O)-, -C(O)CR²R^{2a}-,
 -CR²R^{2a}C(O), -S(O)₂-, -S(O)₂CR²R^{2a}-, -CR²R^{2a}S(O)₂-,
 -NR²S(O)₂-, -S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR²,
 -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

25

Y is a C₃₋₇ monocyclic carbocycle or 3-7 membered monocyclic heterocycle, wherein the carbocycle or heterocycle consists of: carbon atoms and 0-2 heteroatoms selected from N, O, and S(O)_p, the carbocycle or
 30 heterocycle further comprises 0-2 double bonds and 0-2 carbonyl groups, and the carbocycle or heterocycle is substituted with 0-2 R⁴;

alternatively, Y is CY^1Y^2 , and Y^1 and Y^2 are independently C_{1-3} alkyl substituted with 0-1 R^4 ;

5 Z is selected from a bond, CH_2 , CH_2CH_2 , CH_2O , OCH_2 , $C(O)$, NH , CH_2NH , $NHCH_2$, $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $NHC(O)NH$, $NHC(O)CH_2C(O)NH$, $NHC(O)C(O)NH$, $C(O)NHS(O)_2$, $S(O)_2$, $CH_2S(O)_2$, $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$, provided that Z does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;

10 Z^2 is selected from H, C_{1-4} alkyl, phenyl, benzyl, $C(O)R^{3b}$, $S(O)R^{3f}$, and $S(O)_2R^{3f}$;

15 R^{1a} , at each occurrence, is selected from H, $-(CH_2)_r-R^{1b}$, $-(CH(CH_3))_r-R^{1b}$, $-(C(CH_3)_2)_r-R^{1b}$, $-O-(CR^3R^{3a})_r-R^{1b}$, $-NR^2-(CR^3R^{3a})_r-R^{1b}$, and $-S-(CR^3R^{3a})_r-R^{1b}$, provided that R^{1a} forms other than an N-halo, N-S, O-O, or N-CN bond;

20 alternatively, when two R^{1a} groups are attached to adjacent atoms or to the same carbon atom, together with the atoms to which they are attached they form a 5-7 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b} and 0-3 ring double bonds;

25 R^{1b} is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, F, Cl, Br, I, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$, $NR^2C(O)NHR^2$, $NR^2C(O)_2R^{2a}$, $OC(O)NR^2R^{2a}$, $C(O)NR^2R^{2a}$, $C(O)NR^2(CH_2)_rOR^2$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle consisting of carbon atoms and from 1-4 heteroatoms

selected from the group consisting of N, O, and S(O)_p and substituted with 0-2 R^{4b}, provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

5 R², at each occurrence, is selected from H, CF₃, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₅₋₆ carbocycle substituted with 0-2 R^{4b}, a C₅₋₆ carbocycle-CH₂- substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon
10 atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2a}, at each occurrence, is selected from H, CF₃, CH₃,
15 CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group
20 consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated
25 ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2b}, at each occurrence, is selected from CF₃, C₁₋₄ alkoxy,
30 C₁₋₆ alkyl substituted with 0-3 R^{4b}, benzyl, C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 4-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

- R^{2c} , at each occurrence, is selected from CF_3 , OH, C_{1-4} alkoxy, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 10 R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the
- 15 group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p-S(O)_p$, S-O, O-N, O-S, or O-O moiety;
- 20 alternatively, when two R^{2d} 's are attached to the same nitrogen atom, then R^{2d} and R^{2d} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and
- 25 consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the
- 30 group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a C(O)-halo or C(O)- $S(O)_p$ moiety;

R^3 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

- 5 R^{3a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

alternatively, R^3 and R^{3a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6
10 membered saturated, partially unsaturated, or unsaturated ring consisting of: carbon atoms and the nitrogen atom to which R^3 and R^{3a} are attached;

- R^{3c} , at each occurrence, is selected from CH_3 , CH_2CH_3 ,
15 $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, CH_2 -phenyl, CH_2CH_2 -phenyl, and $C(=O)R^{3c}$;

20

R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, cyclopropyl-methyl, benzyl, and phenyl;

- 25 alternatively, when R^3 and R^{3g} are attached to the same carbon atom, they combine with the attached carbon atom to form a cyclopropyl group;

R^4 , at each occurrence, is selected from H, $=O$, OR^2 , CH_2OR^2 ,
30 $(CH_2)_2OR^2$, F, Cl, Br, I, C_{1-4} alkyl, $-CN$, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $(CH_2)_2NR^2R^{2a}$, $C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , 5-6 membered carbocycle substituted with 0-1 R^5 , and a 5-6 membered heterocycle substituted with 0-1 R^5 and consisting of:

carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

5 R^{4b}, at each occurrence, is selected from H, =O, OR³,
 CH₂OR³, F, Cl, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂,
 CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, -CN,
 NO₂, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, CH₂-C(O)R³, C(O)OR^{3c},
 CH₂C(O)OR^{3c}, NR³C(O)R^{3a}, CH₂NR³C(O)R^{3a}, C(O)NR³R^{3a},
 CH₂C(O)NR³R^{3a}, NR³C(O)NR³R^{3a}, CH₂NR³C(O)NR³R^{3a},
 10 C(=NR³)NR³R^{3a}, CH₂C(=NR³)NR³R^{3a}, NR³C(=NR³)NR³R^{3a},
 CH₂NR³C(=NR³)NR³R^{3a}, SO₂NR³R^{3a}, CH₂SO₂NR³R^{3a},
 NR³SO₂NR³R^{3a}, CH₂NR³SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl,
 CH₂NR³SO₂-C₁₋₄ alkyl, NR³SO₂CF₃, CH₂NR³SO₂CF₃,
 NR³SO₂-phenyl, CH₂NR³SO₂-phenyl, S(O)_pCF₃, CH₂S(O)_pCF₃,
 15 S(O)_p-C₁₋₄ alkyl, CH₂S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl,
 CH₂S(O)_p-phenyl, CF₃, and CH₂-CF₃;

R^{4c}, at each occurrence, is selected from =O, (CR³R^{3a})_rOR²,
 (CR³R^{3a})_rF, (CR³R^{3a})_rBr, (CR³R^{3a})_rCl, (CR³R^{3a})_rCF₃, C₁₋₄
 20 alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, (CR³R^{3a})_rCN,
 (CR³R^{3a})_rNO₂, (CR³R^{3a})_rNR²R^{2a}, (CR³R^{3a})_rN(→O)R²R^{2a},
 (CR³R^{3a})_rC(O)R^{2c}, (CR³R^{3a})_rNR²C(O)R^{2b},
 (CR³R^{3a})_rC(O)NR²R^{2a}, (CR³R^{3a})_rNR²C(O)NR²R^{2a},
 (CR³R^{3a})_rSO₂NR²R^{2a}, (CR³R^{3a})_rNR²SO₂NR²R^{2a},
 25 (CR³R^{3a})_rNR²SO₂R^{5a}, (CR³R^{3a})_rS(O)_pR^{5a}, (CF₂)_rCF₃,
 (CR³R^{3a})_rC₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and
 (CR³R^{3a})_r5-10 membered heterocycle substituted with 0-2
 R^{4b} and consisting of carbon atoms and from 1-4
 heteroatoms selected from the group consisting of N,
 30 O, and S(O)_p;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$,
5 $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $NR^3C(O)NR^3R^{3a}$,
 $CH(=NOR^{3d})$, $C(=NR^3)NR^3R^{3a}$, $NR^3C(=NR^3)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 $NR^3SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -
phenyl, $S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 ,
phenyl substituted with 0-2 R^6 , naphthyl substituted
10 with 0-2 R^6 , and benzyl substituted with 0-2 R^6 ;

R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$,
 $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, -CN, NO_2 , NR^2R^{2a} ,
15 $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$,
 $NR^2C(O)NR^2R^{2a}$, $C(=NH)NH_2$, $NHC(=NH)NH_2$, $SO_2NR^2R^{2a}$,
 $NR^2SO_2NR^2R^{2a}$, and $NR^2SO_2C_{1-4}$ alkyl; and,

r , at each occurrence, is selected from 0, 1, 2, and 3.
20

[10] In another preferred embodiment, the present invention provides a novel compound, wherein:

25 ring M, including M_1 , M_2 , and, if present, M_3 , is selected
from phenyl, pyrrole, furan, thiophene, pyrazole,
imidazole, isoxazole, oxazole, isothiazole, thiazole,
1,2,3-triazole, 1,2,4-triazole, 1,3,4-triazole, 1,2,3-
oxadiazole, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,3-
30 thiadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole,
1,2,3,4-tetrazole, 1,2,3,5-tetrazole, pyran,
thiopyran, thiopyran-1,1-dioxide, pyridine,
pyrimidine, pyridazine, pyrazine, 1,2,3-triazine,
1,2,4-triazine, 1,2,3,4-tetrazine, dihydro-pyrrole,

dihydro-furan, dihydro-thiophene, dihydro-pyrazole,
dihydro-imidazole, dihydro-isoxazole, dihydro-oxazole,
dihydro-isothiazole, dihydro-thiazole, dihydro-1,2,3-
triazole, dihydro-1,2,4-triazole, dihydro-1,3,4-
5 triazole, dihydro-1,2,3-oxadiazole, dihydro-1,2,4-
oxadiazole, dihydro-1,3,4-oxadiazole, dihydro-1,2,3-
thiadiazole, dihydro-1,2,4-thiadiazole, dihydro-1,3,4-
thiadiazole, dihydro-1,2,3,4-tetrazole, dihydro-
1,2,3,5-tetrazole, dihydro-pyran, dihydro-thiopyran,
10 dihydro-thiopyran-1,1-dioxide, dihydro-pyridine,
dihydro-pyrimidine, dihydro-pyridazine, dihydro-
pyrazine, dihydro-1,2,3-triazine, dihydro-1,2,4-
triazine, dihydro-1,2,3,4-tetrazine, cyclopropane,
cyclobutane, cyclopentene, cyclopentane, cyclohexene,
15 cyclohexane, cycloheptane, tetrahydro-pyrrole,
tetrahydro-furan, tetrahydro-thiophene, tetrahydro-
thiophene-1,1-dioxide, tetrahydro-pyrazole,
tetrahydro-imidazole, tetrahydro-isoxazole,
tetrahydro-oxazole, tetrahydro-isothiazole,
20 tetrahydro-thiazole, tetrahydro-1,2,3-triazole,
tetrahydro-1,2,4-triazole, tetrahydro-1,3,4-triazole,
tetrahydro-1,2,3-oxadiazole, tetrahydro-1,2,4-
oxadiazole, tetrahydro-1,3,4-oxadiazole, tetrahydro-
1,2,3-thiadiazole, tetrahydro-1,2,4-thiadiazole,
25 tetrahydro-1,3,4-thiadiazole, tetrahydro-1,2,3,4-
tetrazole, tetrahydro-1,2,3,5-tetrazole, tetrahydro-
pyran, tetrahydro-thiopyran, tetrahydro-thiopyran-1,1-
dioxide, tetrahydro-pyridine, tetrahydro-pyrimidine,
tetrahydro-pyridazine, tetrahydro-pyrazine,
30 tetrahydro-1,2,3-triazine, tetrahydro-1,2,4-triazine,
tetrahydro-1,2,3,4-tetrazine, piperidine, indan,
isothiazolidine 1,1-dioxide, [1,2]thiazinane 1,1-
dioxide, 1,2,3,4-tetrahydro-naphthalene, 7,8-dimethyl-
1-oxa-spiro[4.4]nonane, 6,7-dihydro-5H-[1]pyrindine,
35 6,7-dihydro-5H-[2]pyrindine, 5,6,7,8-tetrahydro-
quinoline, 5,6,7,8-tetrahydro-isoquinoline, 5,6,7,8-

tetrahydro-quinoxaline, 6,7-dihydro-5H-cyclopentapyrazine, 4,5,6,7-tetrahydro-1H-benzoimidazole, 4,5,6,7-tetrahydro-benzothiazole, 4,5,6,7-tetrahydro-benzooxazole, 4,5,6,7-tetrahydro-5
benzo[c]isothiazole, 4,5,6,7-tetrahydro-benzo[c]isoxazole, 4,5,6,7-tetrahydro-2H-indazole, 4,5,6,7-tetrahydro-2H-isoindole, 4,5,6,7-tetrahydro-1H-indole, 5,6,7,8-tetrahydro-tetrazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-imidazo[1,2-a]pyridine, 10
4,5,6,7-tetrahydro-pyrazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-[1,2,4]triazolo[1,5-a]pyridine, 6,7-dihydro-5H-pyrrolo[1,2-c]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-a]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-b][1,2,4]triazole, 6,7-dihydro-5H-pyrrolotetrazole, 15
5,6-dihydro-4H-pyrrolo[1,2-b]pyrazole, 5,6-dihydro-4H-cyclopenta[d]isoxazole, 5,6-dihydro-4H-cyclopentaoxazole, 5,6-dihydro-4H-cyclopenta[c]isoxazole, 5,6-dihydro-4H-cyclopenta[d]isothiazole, 5,6-dihydro-4H-cyclopentathiazole, 5,6-dihydro-4H-cyclopenta[c]isothiazole, 1,4,5,6-tetrahydro-cyclopentapyrazole, 1,4,5,6-tetrahydro-cyclopentaimidazole, 2,4,5,6-tetrahydro-cyclopentapyrazole, 5,6-dihydro-4H-cyclopenta[1,2,5]thiadiazole, 5,6-dihydro-4H-cyclopenta[1,2,5]oxadiazole, 5,6-dihydro-4H-cyclopenta[c]furan, 2,4,5,6-tetrahydro-cyclopenta[c]pyrrole, 5,6-dihydro-4H-cyclopenta[b]furan, 5,6-dihydro-4H-cyclopenta[c]thiophene, 5,6-dihydro-4H-cyclopenta[b]furan, 5,6-dihydro-4H-cyclopenta[b]thiophene, 1,4,5,6-tetrahydro-cyclopenta[b]pyrrole, 2,3-dihydro-1H-indolizin-5-one, 6,7,8,9-tetrahydro-quinolizin-4-one, 1-oxa-35
spiro[4.4]nonane, 1-aza-spiro[4.4]nonane, 2-oxa-spiro[4.4]nonane, 2-aza-spiro[4.4]nonane, 1-aza-

spiro[4.5]decane, 1-oxa-spiro[4.5]decane, 2-oxa-spiro[4.5]decane, 2-aza-spiro[4.5]decane, 1-thia-spiro[4.4]nonane, 1-thia-spiro[4.5]decane, 2-thia-spiro[4.4]nonane, 2-thia-spiro[4.5]decane, 7-oxa-bicyclo[2.2.1]heptane, 2-oxa-bicyclo[2.2.1]heptane, 7-thia-bicyclo[2.2.1]heptane, 2-thia-bicyclo[2.2.1]heptane, 2-aza-bicyclo[2.2.1]heptane, 7-aza-bicyclo[2.2.1]heptane, 4,5,6,7-tetrahydro-benzo[d]isoxazole, 4,5,6,7-tetrahydro-benzooxazole, 4,5,6,7-tetrahydro-benzo[d]isothiazole, 4,5,6,7-tetrahydro-benzothiazole, 4,5,6,7-tetrahydro-1H-indazole, 4,5,6,7-tetrahydro-benzo[c]thiophene, 4,5,6,7-tetrahydro-benzo[b]thiophene, 4,5,6,7-tetrahydro-isobenzofuran, 4,5,6,7-tetrahydro-benzofuran, 5,6,7,8-tetrahydro-quinoxaline, 6,7-dihydro-5H-cyclopentapyrazine, 5,6,7,8-tetrahydro-imidazo[1,5-a]pyridine, 5,6,7,8-tetrahydro-imidazo[1,2-a]pyridine, 5,6,7,8-tetrahydro-[1,2,4]triazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-tetrazolo[1,5-a]pyridine, 4,5,6,7-tetrahydro-pyrazolo[1,5-a]pyridine, 6,7-dihydro-5H-pyrrolo[1,2-a]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-b][1,2,4]triazole, 5,6-dihydro-4H-pyrrolo[1,2-b]pyrazole, and 6,7-dihydro-5H-pyrrolotetrazole;

25

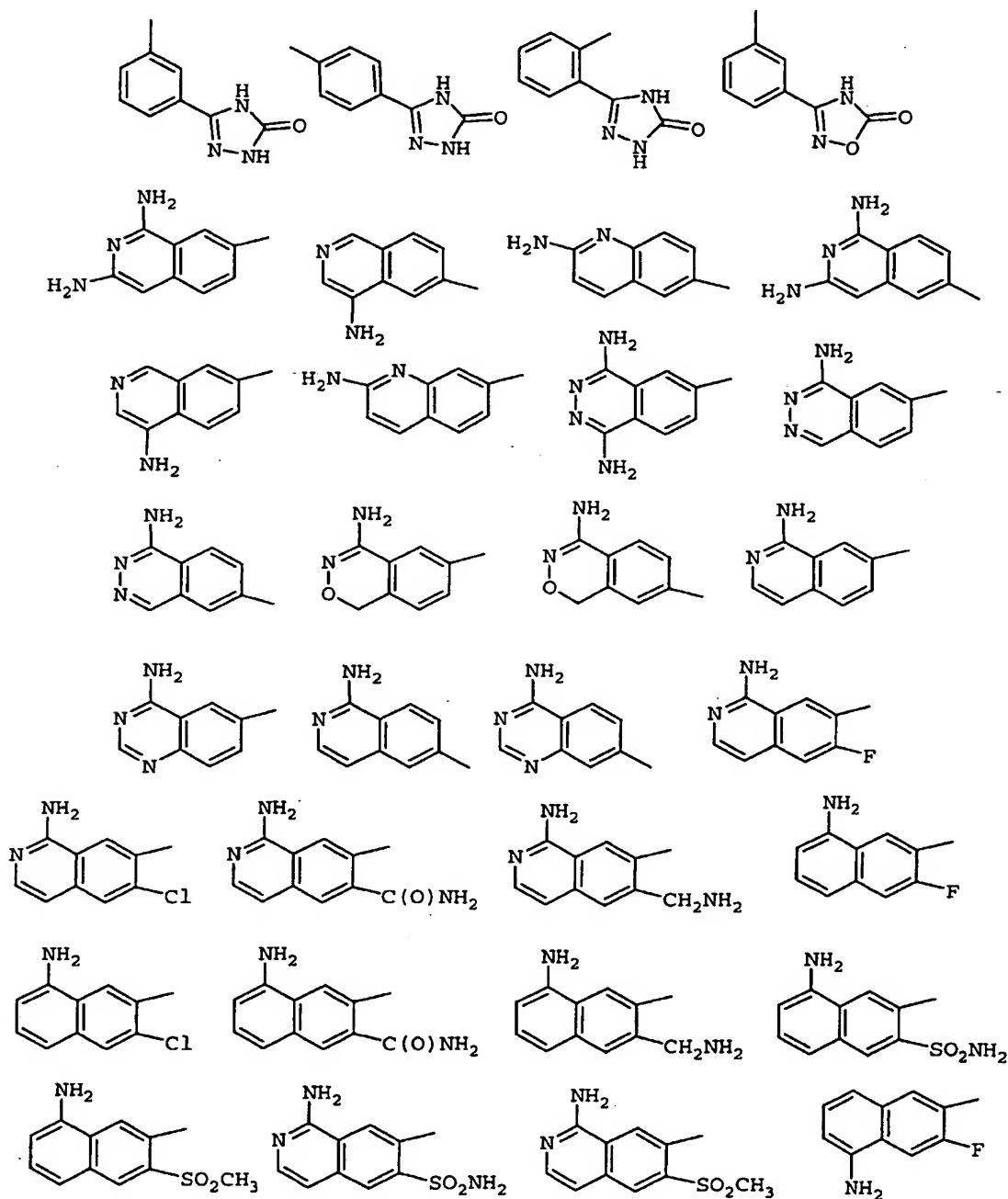
ring M is substituted with 0-3 R^{1a} and 0-1 carbonyl group;

G is selected from the group:

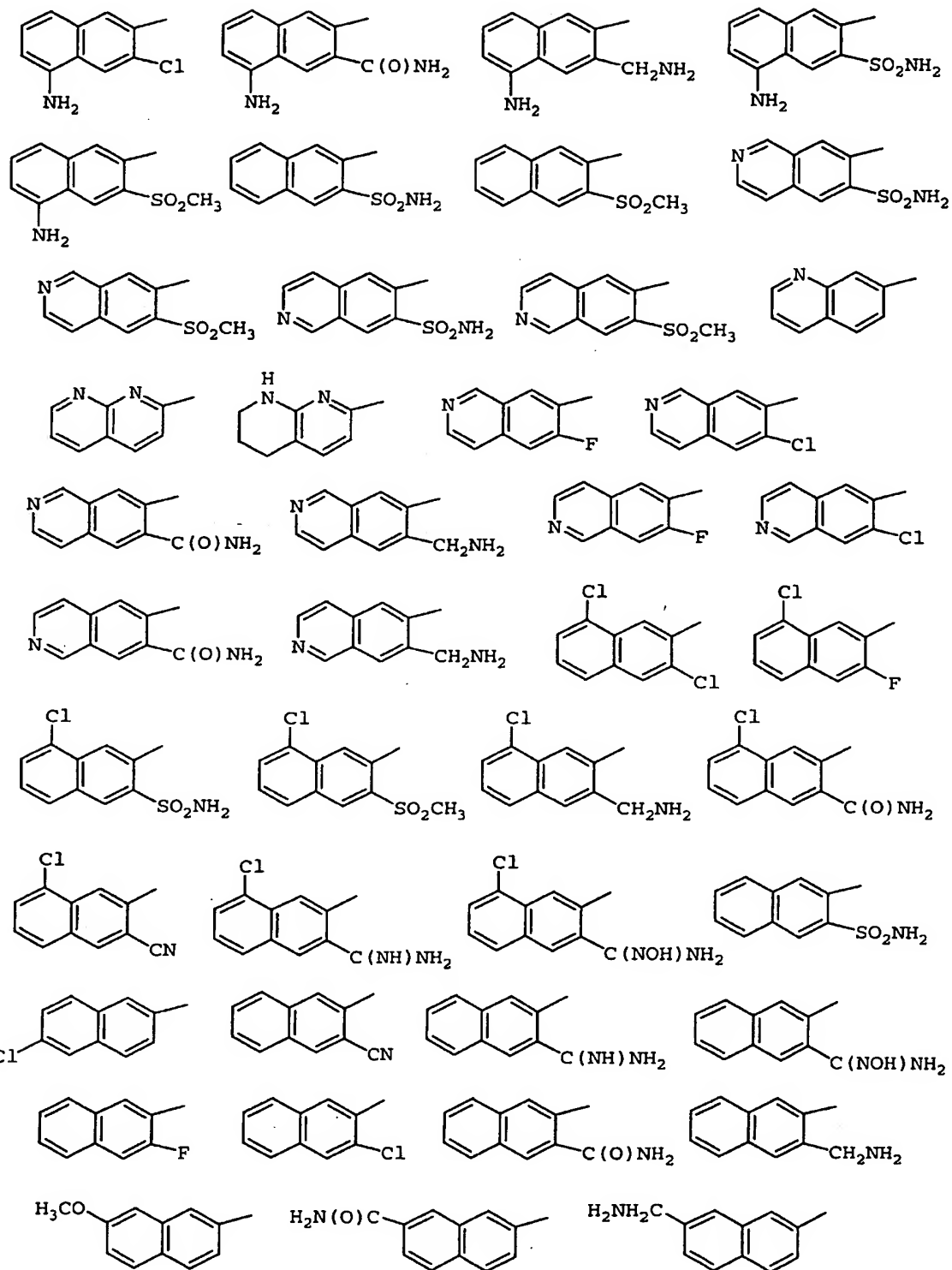
phenyl; 4-ethyl-phenyl; 2,5-bis-aminomethyl-phenyl;
30 2-amido-4-methoxy-phenyl; 2-amido-5-chloro-phenyl;
2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
2-aminomethyl-3-methoxy-phenyl;
2-aminomethyl-4-fluoro-phenyl;
2-aminomethyl-4-methoxy-phenyl;
35 2-aminomethyl-5-fluoro-phenyl;
2-aminomethyl-5-methoxy-phenyl;

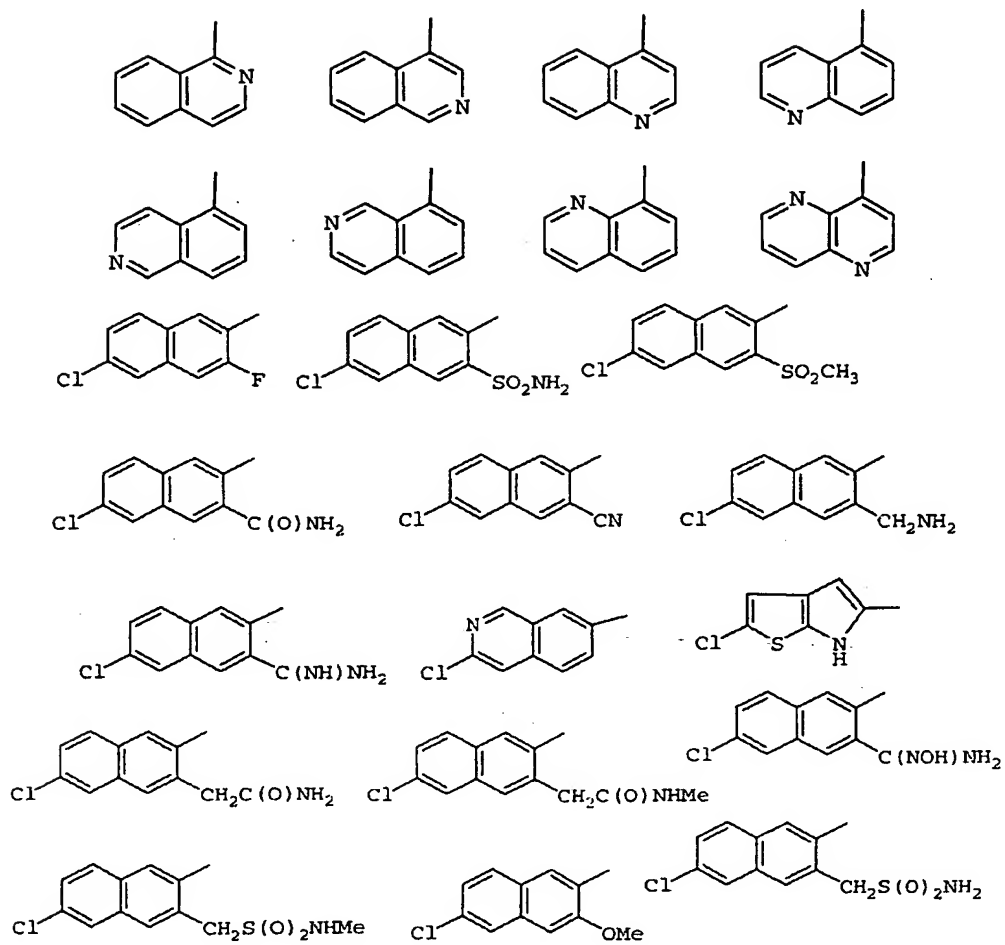
- 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
2-aminosulfonyl-phenyl; 2-hydroxy-4-methoxy-phenyl;
2-methylsulfonyl-phenyl;
- 5 3-(N,N-dimethylamino)-4-chloro-phenyl;
3-(N,N-dimethylamino)-phenyl; 3-(N-hydroxy-amidino)-phenyl;
3-(N-methoxy-amidino)-phenyl;
3-(N-methylamino)-4-chloro-phenyl;
3-(N-methylamino)-phenyl; 3-amidino-phenyl;
- 10 3-amido-6-hydroxy-phenyl; 3-amido-phenyl;
3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
3-amino-phenyl; 3-chloro-4-fluoro-phenyl; 3-chloro-phenyl;
3-hydroxy-4-methoxy-phenyl; 3,5-dichloro-thien-2-yl;
4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
- 15 4-(N-methylamino)-5-chloro-thien-2-yl;
4-amino-5-chloro-thien-2-yl; 4-amino-pyrid-2-yl;
4-chloro-3-fluoro-phenyl; 4-chloro-phenyl;
4-chloro-pyrid-2-yl; 4-methoxy-2-methylsulfonyl-phenyl;
4-methoxy-phenyl; 2-methoxy-pyrid-5-yl;
- 20 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
5-(N-methylamino)-4-chloro-thien-2-yl;
5-amino-4-chloro-thien-2-yl;
5-chloro-2-aminosulfonyl-phenyl;
5-chloro-2-methylsulfonyl-phenyl; 5-chloro-pyrid-2-yl;
- 25 5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-chloro-
pyrimidin-3-yl; 6-chloro-pyridazin-3-yl;
2-aminomethyl-4-chloro-phenyl;
2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl;
- 30 4-chloro-2-methylsulfonyl-phenyl;
2-aminosulfonyl-4-fluoro-phenyl; 2-amido-4-fluoro-phenyl;
4-fluoro-2-methylsulfonyl-phenyl;
2-aminomethyl-4-bromo-phenyl;
2-aminosulfonyl-4-bromo-phenyl; 2-amido-4-bromo-phenyl;
- 35 4-bromo-2-methylsulfonyl-phenyl;
2-aminomethyl-4-methyl-phenyl;

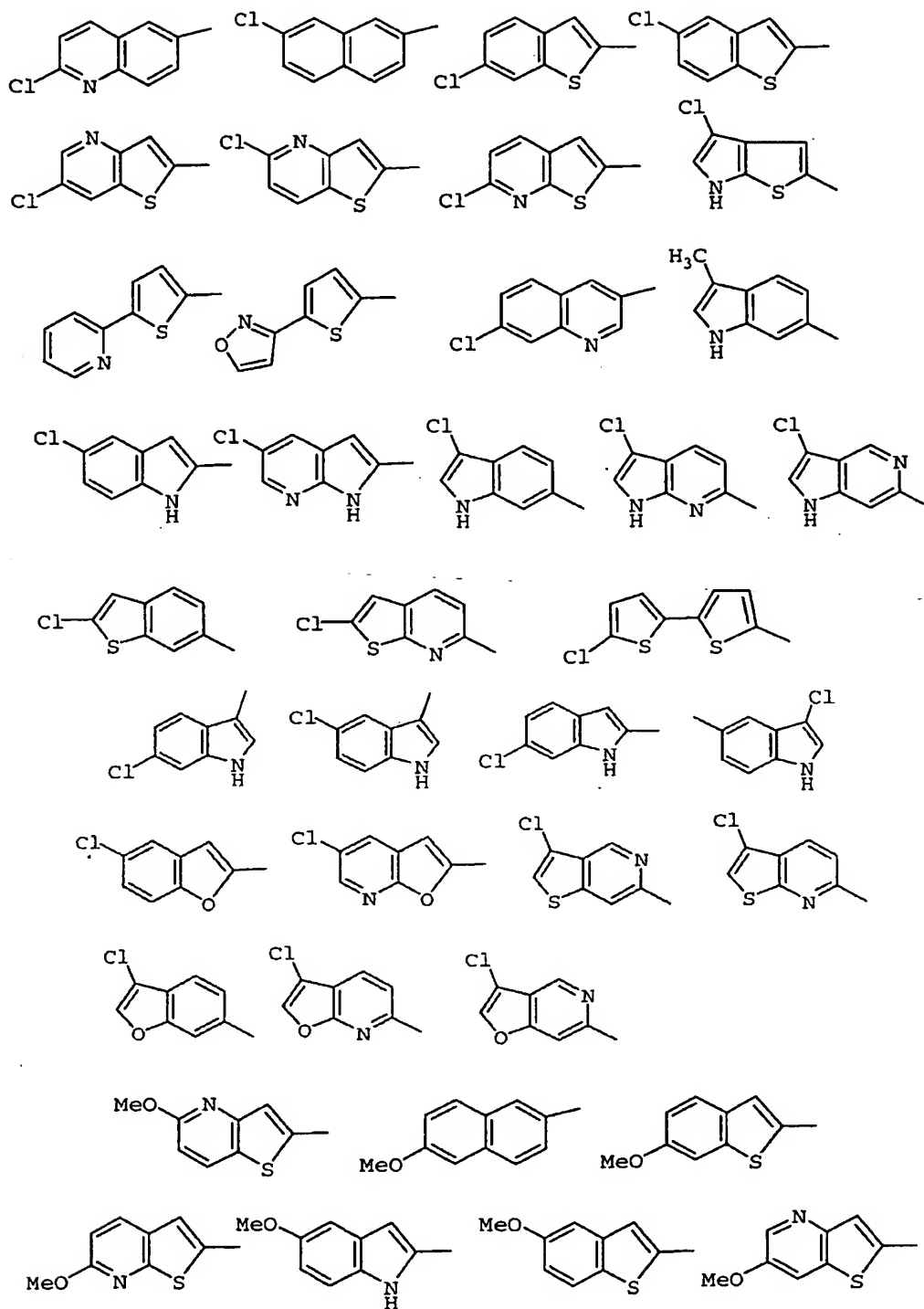
- 2-aminosulfonyl-4-methyl-phenyl; 2-amido-4-methyl-phenyl;
 2-methylsulfonyl-4-methyl-phenyl; 4-fluoro-pyrid-2-yl;
 4-bromo-pyrid-2-yl; 4-methyl-pyrid-2-yl;
 5-fluoro-thien-2-yl; 5-bromo-thien-2-yl;
 5 5-methyl-thien-2-yl; 2-amido-4-methoxy-phenyl;

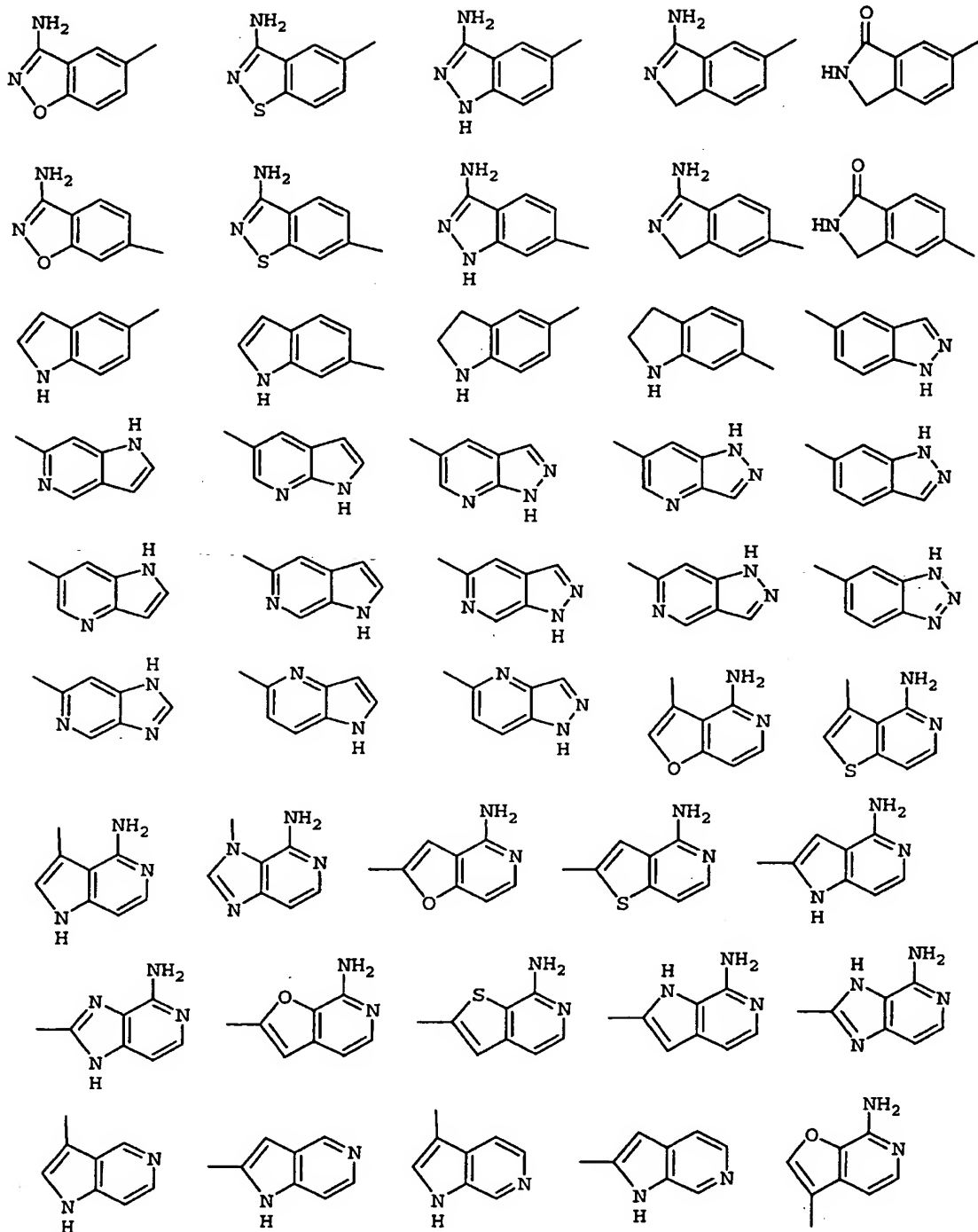


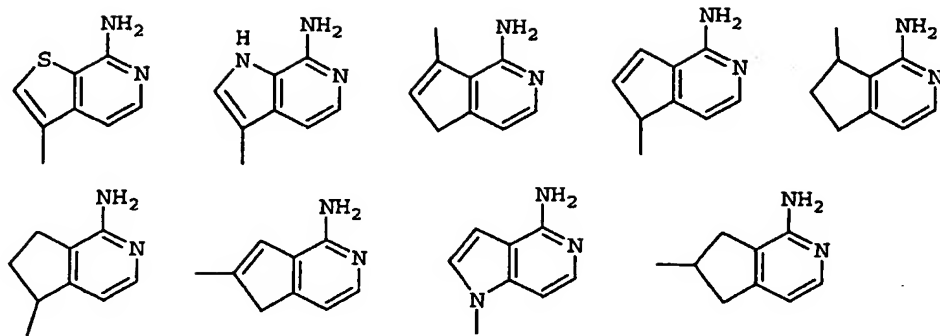
10











G_1 is absent or is selected from $(CR^3R^{3a})_{1-3}$, $CR^3=CR^3$,
 $(CR^3R^{3a})_u C(O)(CR^3R^{3a})_w$, $(CR^3R^{3a})_u O(CR^3R^{3a})_w$,
 5 $(CR^3R^{3a})_u NR^{3b}(CR^3R^{3a})_w$, $(CR^3R^{3a})_u C(O)NR^{3b}(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u NR^{3b}C(O)(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u NR^{3b}C(O)(CR^3R^{3a})_u C(O)NR^{3b}(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u S(CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O)(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u S(O)_2(CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O)NR^{3b}(CR^3R^{3a})_w$,
 10 $(CR^3R^{3a})_u NR^{3b}S(O)_2(CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O)_2NR^{3b}(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u C(O)NR^{3b}S(O)_2(CR^3R^{3a})_w$,
 $(CR^3R^{3a})_u NR^{3b}C(S)(CR^3R^{3a})_u C(O)NR^{3b}(CR^3R^{3a})_w$, and
 $(CR^3R^{3a})_u NR^{3b}C(O)(CR^3R^{3a})_u C(S)NR^{3b}(CR^3R^{3a})_w$, wherein u
 $+ w$ total 0, 1, or 2, provided that G_1 does not form a
 15 N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to
 which it is attached;

A is selected from one of the following carbocycles and
 heterocycles which are substituted with 0-2 R^4 ;
 20 cyclohexyl, phenyl, piperidinyl, piperazinyl,
 pyridyl, pyrimidyl, furanyl, morpholinyl, thienyl,
 pyrrolyl, pyrrolidinyl, oxazolyl, isoxazolyl,
 thiazolyl, isothiazolyl, pyrazolyl, imidazolyl,
 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl,
 25 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl,
 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl,
 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl,

1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl,
1,3,4-triazolyl, benzofuranyl, benzothiofuranyl,
indoliny, indolyl, benzimidazolyl, benzoxazolyl,
benzthiazolyl, indazolyl, benzisoxazolyl,
5 benzisothiazolyl, and isoindazolyl;

X is selected from $-(CR^2R^{2a})_{1-2}-$, $-C(O)-$, $-S(O)_2-$,
 $-NR^2S(O)_2-$, $-NR^2S(O)_2NR^2-$, $-NR^2C(O)-$, $-C(O)NR^2-$, NR^2 ,
 $-NR^2CR^2R^{2a}-$, $-CR^2R^{2a}NR^2-$, O, $-OCR^2R^{2a}-$, and $-CR^2R^{2a}O-$;

10

Y is a C_{3-6} monocyclic carbocycle or 5-6 membered monocyclic
heterocycle, wherein the carbocycle or heterocycle
consists of carbon atoms and 0-2 heteroatoms selected
from N, O, and $S(O)_p$, the carbocycle or heterocycle
15 further comprises 0-1 double bonds and 0-1 carbonyl
groups, and the carbocycle or heterocycle is
substituted with 0-2 R^4 ;

alternatively, Y is CY^1Y^2 , and Y^1 and Y^2 are independently
20 C_{1-2} alkyl substituted with 0-1 R^4 ;

R^{1a} , at each occurrence, is selected from H, R^{1b} ,
 $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, CH_2R^{1b} , and $CH_2CH_2R^{1b}$, provided
that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

25

alternatively, when two R^{1a} groups are attached to adjacent
atoms or to the same carbon atom, together with the
atoms to which they are attached, they form a 5-6
membered ring consisting of: carbon atoms and 0-2
heteroatoms selected from the group consisting of N,
30 O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b}
and comprising: 0-3 double bonds;

R^{1b} is selected from H, CH_3 , CH_2CH_3 , F, Cl, Br, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

10

R^2 , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-2 R^{4b} , benzyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

15

R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

20

R^{2b} , at each occurrence, is selected from CF_3 , C_{1-4} alkoxy, C_{1-5} alkyl substituted with 0-3 R^{4b} , benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 4-6 membered substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

25
30

R^{2c} , at each occurrence, is selected from CF_3 , OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$,

CH(CH₃)₂, benzyl, phenyl substituted with 0-2 R^{4b}, and
5-6 membered aromatic heterocycle substituted with 0-2
R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
5 O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom
to which they are attached, combine to form a 3-6
membered saturated, partially saturated or unsaturated
10 ring substituted with 0-2 R^{4b} and consisting of: 0-1
additional heteroatoms selected from the group
consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl
15 substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted
with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with
0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2
R^{4c} and consisting of: carbon atoms and 1-4
heteroatoms selected from the group consisting of N,
20 O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle
substituted with 0-2 R^{4c} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and S(O)_p, provided that R^{2d} forms
other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-
25 S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl
substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted
with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with
30 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2
R^{4c} consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and S(O)_p,
and -(CR³R^{3a})-5-6 membered heterocycle substituted with
0-2 R^{4c} and consisting of: carbon atoms and 1-4

heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

5 R⁴, at each occurrence, is selected from H, (CH₂)₂OR², CH₂OR², OR², F, Cl, Br, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, -CN, NO₂, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, C(O)R^{2c}, NR²C(O)R^{2b}, C(O)NR²R^{2a}, SO₂NR²R^{2a}, CF₃, and
10 CF₂CF₃;

R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d}, (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)R^{2e}, (CR³R^{3g})_r-C(O)R^{2e}, (CR³R^{3g})_r-OC(O)R^{2e}, (CR³R^{3g})_r-C(O)NR^{2d}R^{2d},
20 (CR³R^{3g})_r-C(O)OR^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)OR^{2d}, (CR³R^{3g})_r-SO₂NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}SO₂R^{2d}, and (CR³R^{3g})_r-S(O)_pR^{2d}, provided that S(O)_pR^{2d} forms other than S(O)₂H or S(O)H;

25 R^{4b}, at each occurrence, is selected from H, =O, OR³, CH₂OR³, F, Cl, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, -CN, NO₂, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, CH₂-C(O)R³, C(O)OR^{3c}, CH₂-C(O)OR^{3c}, NR³C(O)R^{3a}, CH₂NR³C(O)R^{3a}, C(O)NR³R^{3a}, CH₂-C(O)NR³R^{3a}, SO₂NR³R^{3a}, CH₂SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, CH₂NR³SO₂-C₁₋₄ alkyl, NR³SO₂-phenyl, CH₂NR³SO₂-phenyl,
30 S(O)_pCF₃, CH₂S(O)_pCF₃, S(O)_p-C₁₋₄ alkyl, CH₂S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl, CH₂S(O)_p-phenyl, and CF₃;

R^{4c} , at each occurrence, is selected from =O, OR^2 ,
 $(CR^3R^{3a})OR^2$, F, $(CR^3R^{3a})F$, Br, $(CR^3R^{3a})Br$, Cl,
 $(CR^3R^{3a})Cl$, CF_3 , $(CR^3R^{3a})CF_3$, C_{1-4} alkyl, C_{2-3} alkenyl,
5 C_{2-3} alkynyl, -CN, $(CR^3R^{3a})CN$, NO_2 , $(CR^3R^{3a})NO_2$, NR^2R^{2a} ,
 $(CR^3R^{3a})NR^2R^{2a}$, $N(\rightarrow O)R^2R^{2a}$, $(CR^3R^{3a})N(\rightarrow O)R^2R^{2a}$, $C(O)R^{2c}$,
 $(CR^3R^{3a})C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $(CR^3R^{3a})NR^2C(O)R^{2b}$,
 $C(O)NR^2R^{2a}$, $(CR^3R^{3a})C(O)NR^2R^{2a}$, $NR^2C(O)NR^2R^{2a}$,
 $(CR^3R^{3a})NR^2C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $(CR^3R^{3a})SO_2NR^2R^{2a}$,
10 $NR^2SO_2NR^2R^{2a}$, $(CR^3R^{3a})NR^2SO_2NR^2R^{2a}$, $NR^2SO_2R^{5a}$,
 $(CR^3R^{3a})NR^2SO_2R^{5a}$, $S(O)_pR^{5a}$, $(CR^3R^{3a})S(O)_pR^{5a}$, CF_3 ,
 CF_2CF_3 , C_{3-10} carbocycle substituted with 0-2 R^{4b} ,
 $(CR^3R^{3a})C_{3-10}$ carbocycle substituted with 0-2 R^{4b} , 5-10
membered heterocycle substituted with 0-2 R^{4b} and
15 consisting of carbon atoms and from 1-4 heteroatoms
selected from the group consisting of N, O, and $S(O)_p$,
and (CR^3R^{3a}) 5-10 membered heterocycle substituted with
0-2 R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
20 O, and $S(O)_p$;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$,
25 $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -phenyl, $S(O)_pCF_3$,
 $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 , phenyl substituted
with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
benzyl substituted with 0-2 R^6 ;

30

R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN, NO_2 , NR^2R^{2a} ,

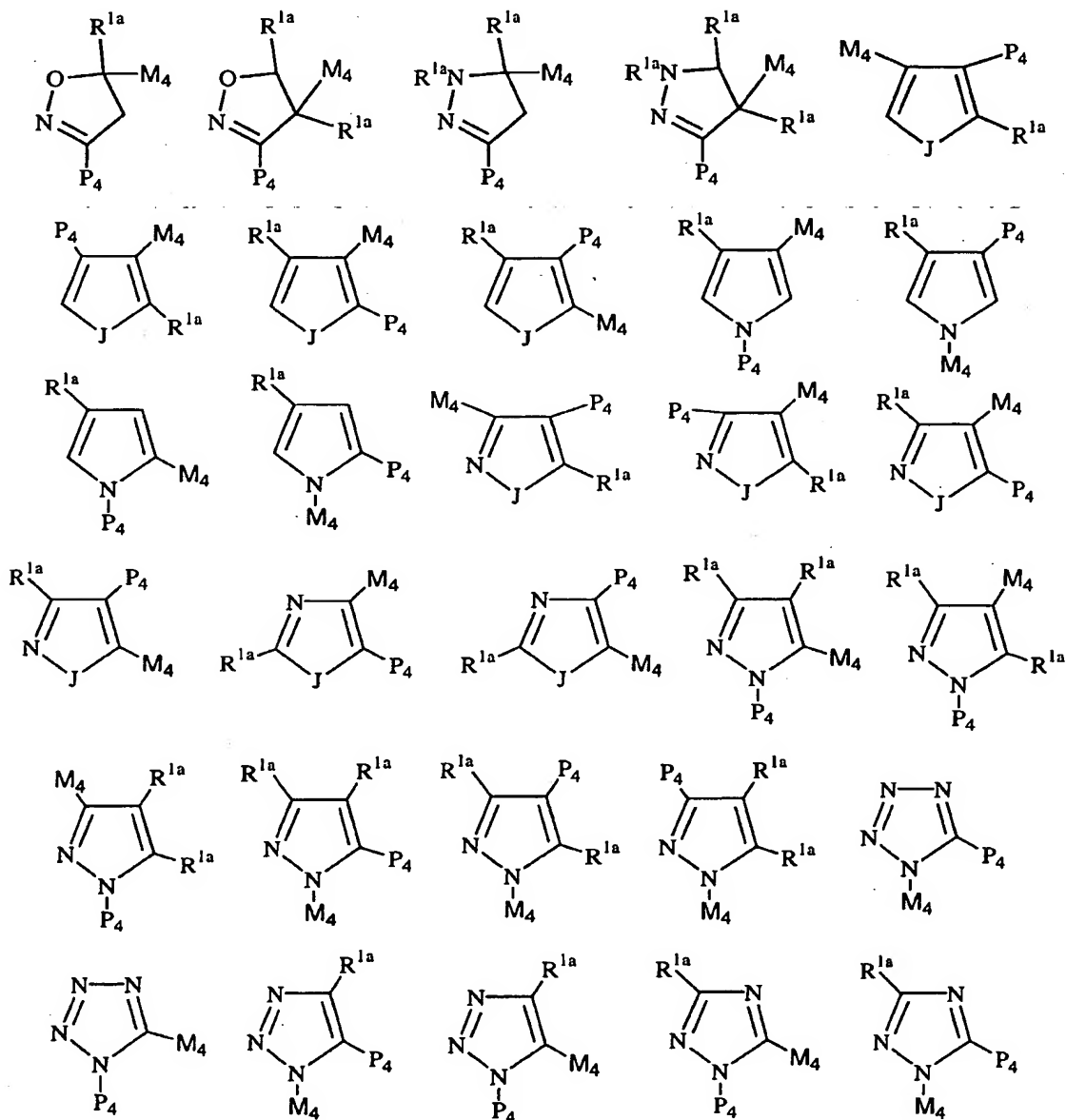
$\text{CH}_2\text{NR}^2\text{R}^{2a}$, C(O)R^{2b} , $\text{CH}_2\text{C(O)R}^{2b}$, $\text{NR}^2\text{C(O)R}^{2b}$, $\text{SO}_2\text{NR}^2\text{R}^{2a}$,
and $\text{NR}^2\text{SO}_2\text{C}_{1-4}$ alkyl; and,

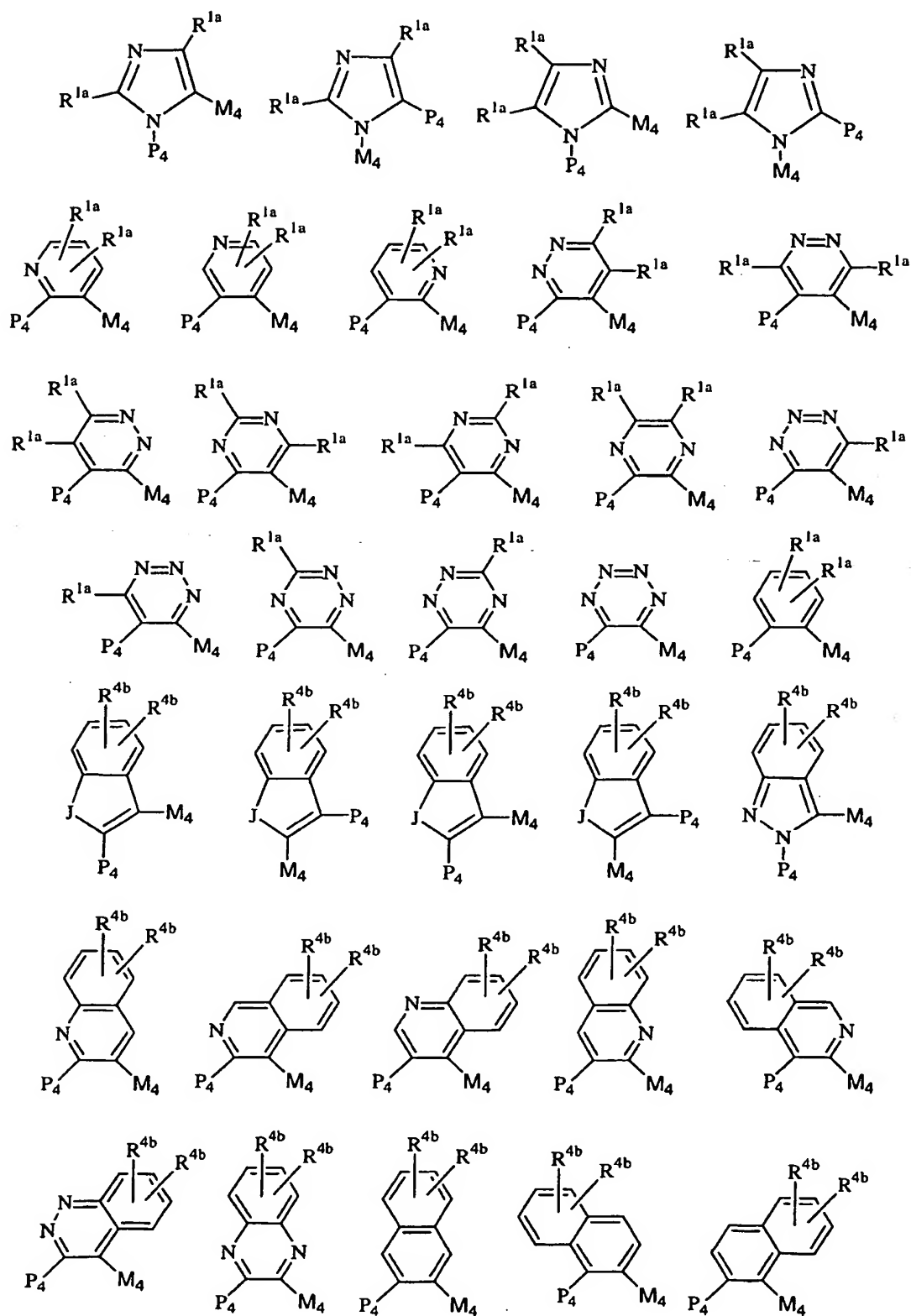
r , at each occurrence, is selected from 0, 1, and 2.

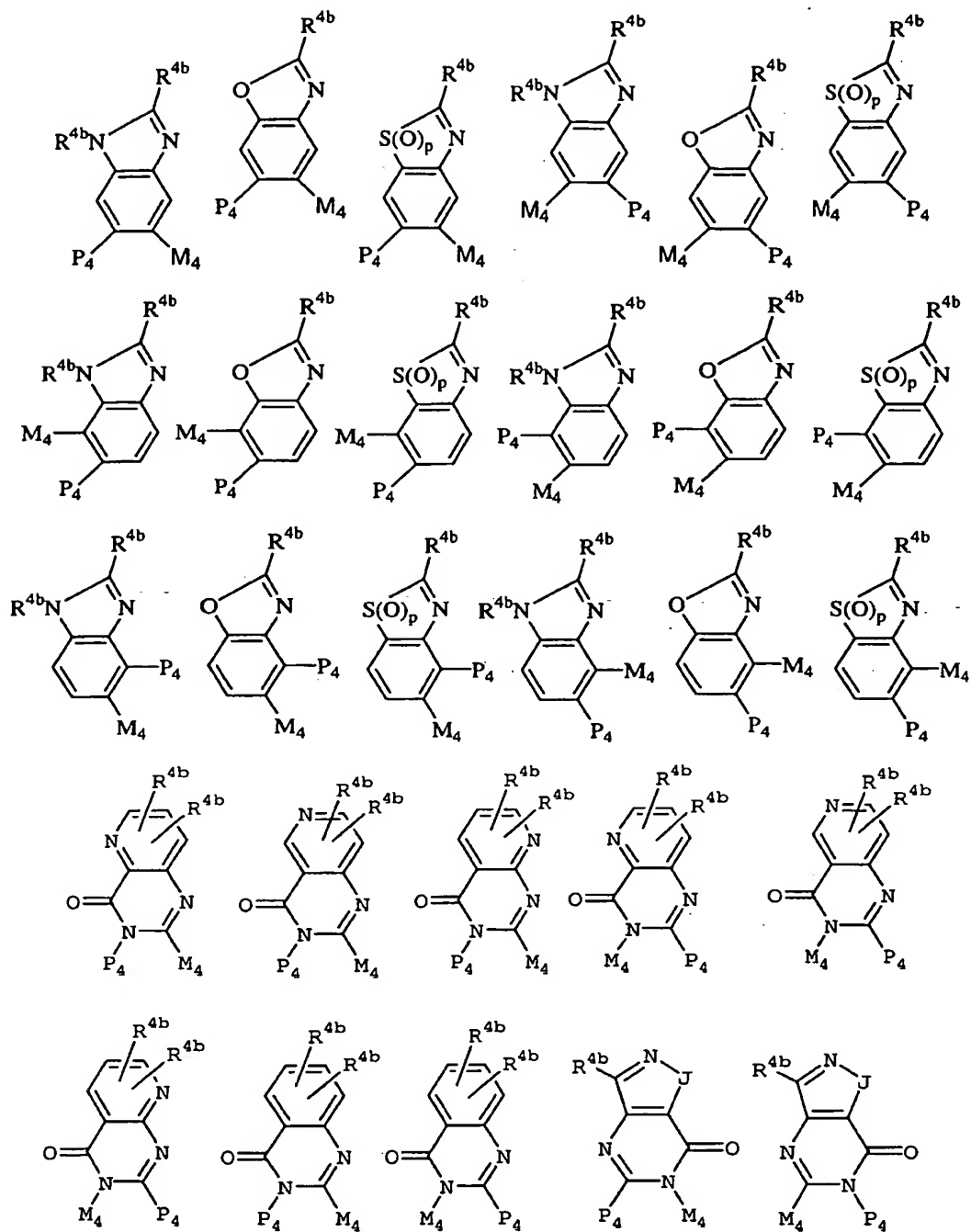
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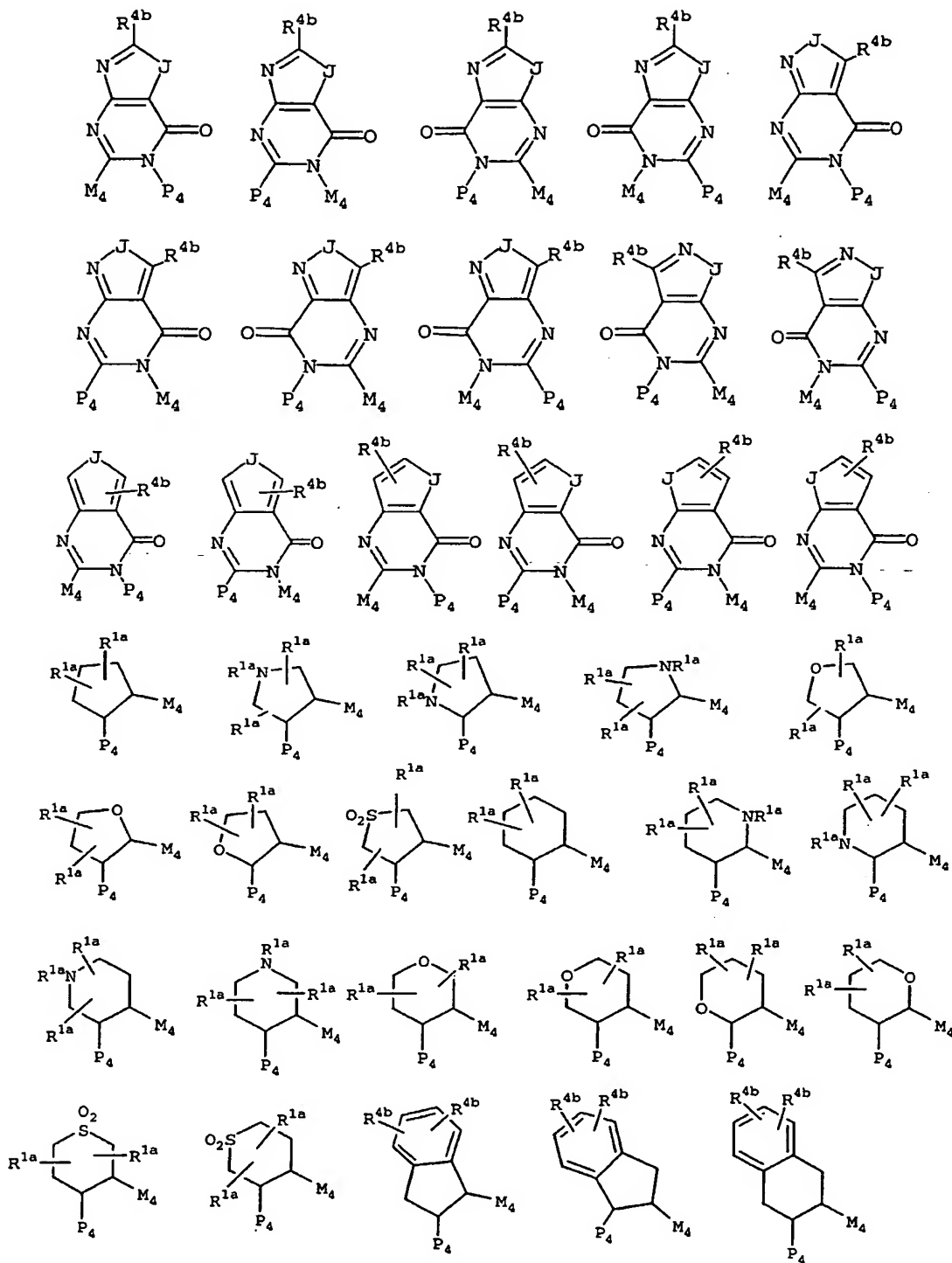
[11] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is selected from:

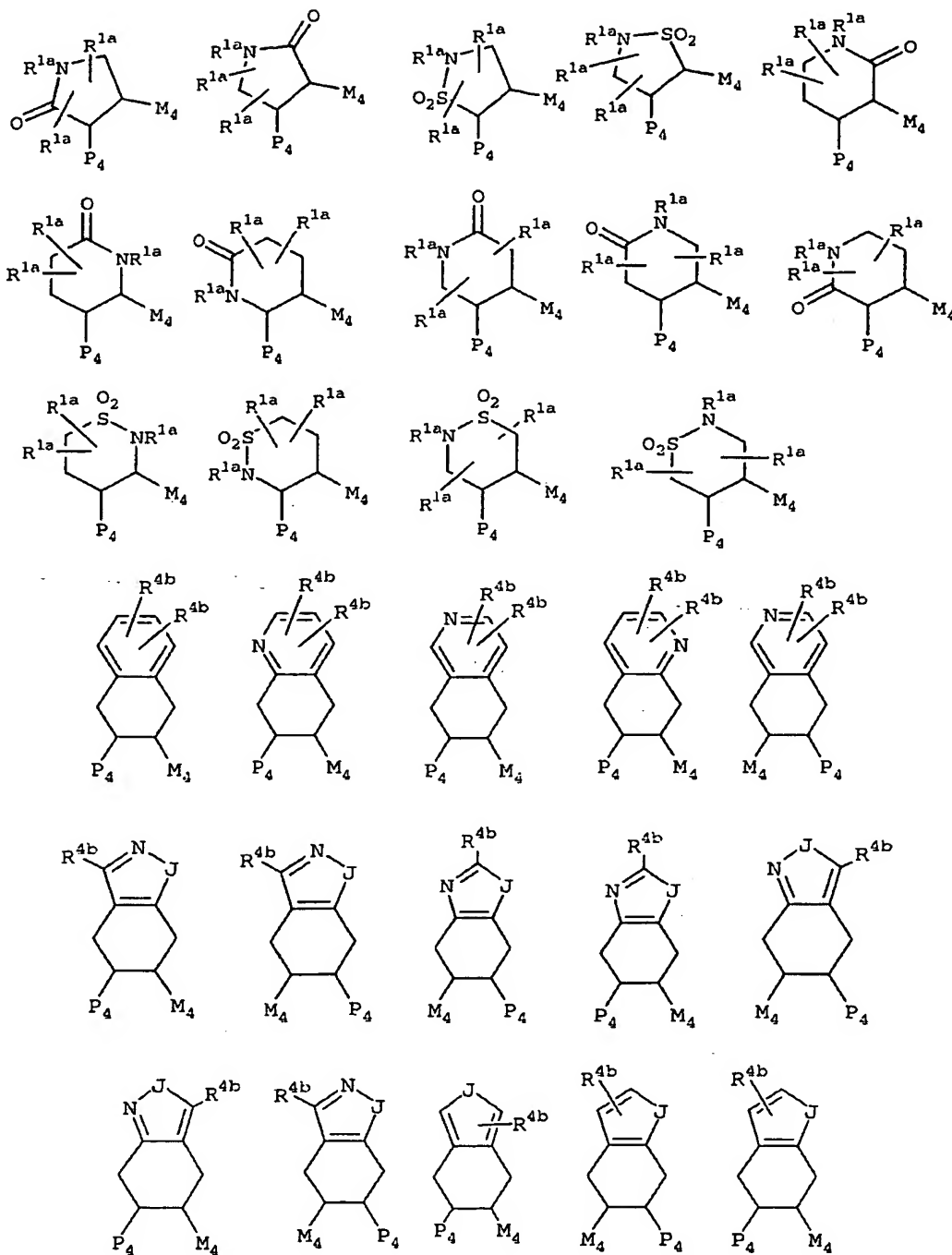
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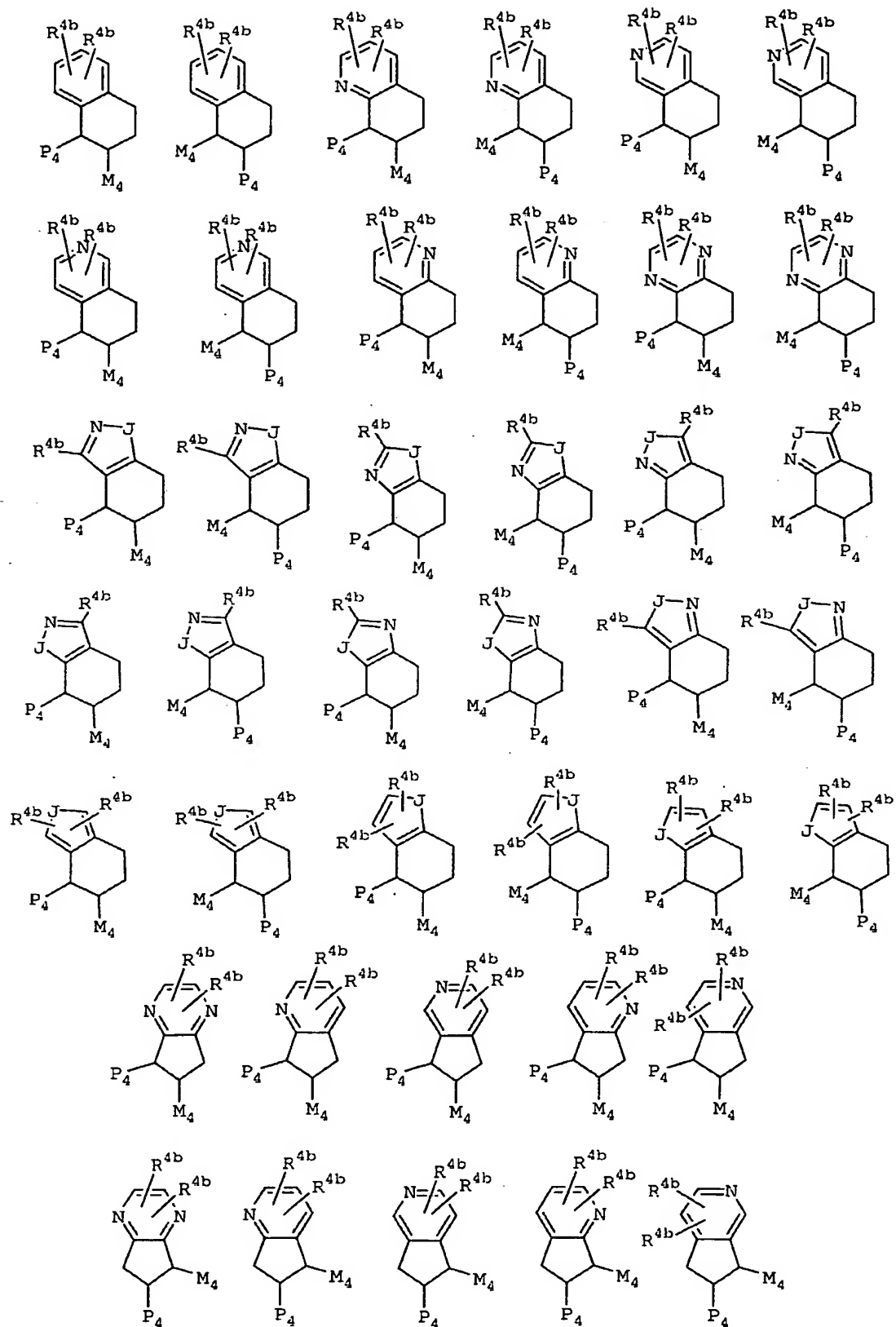


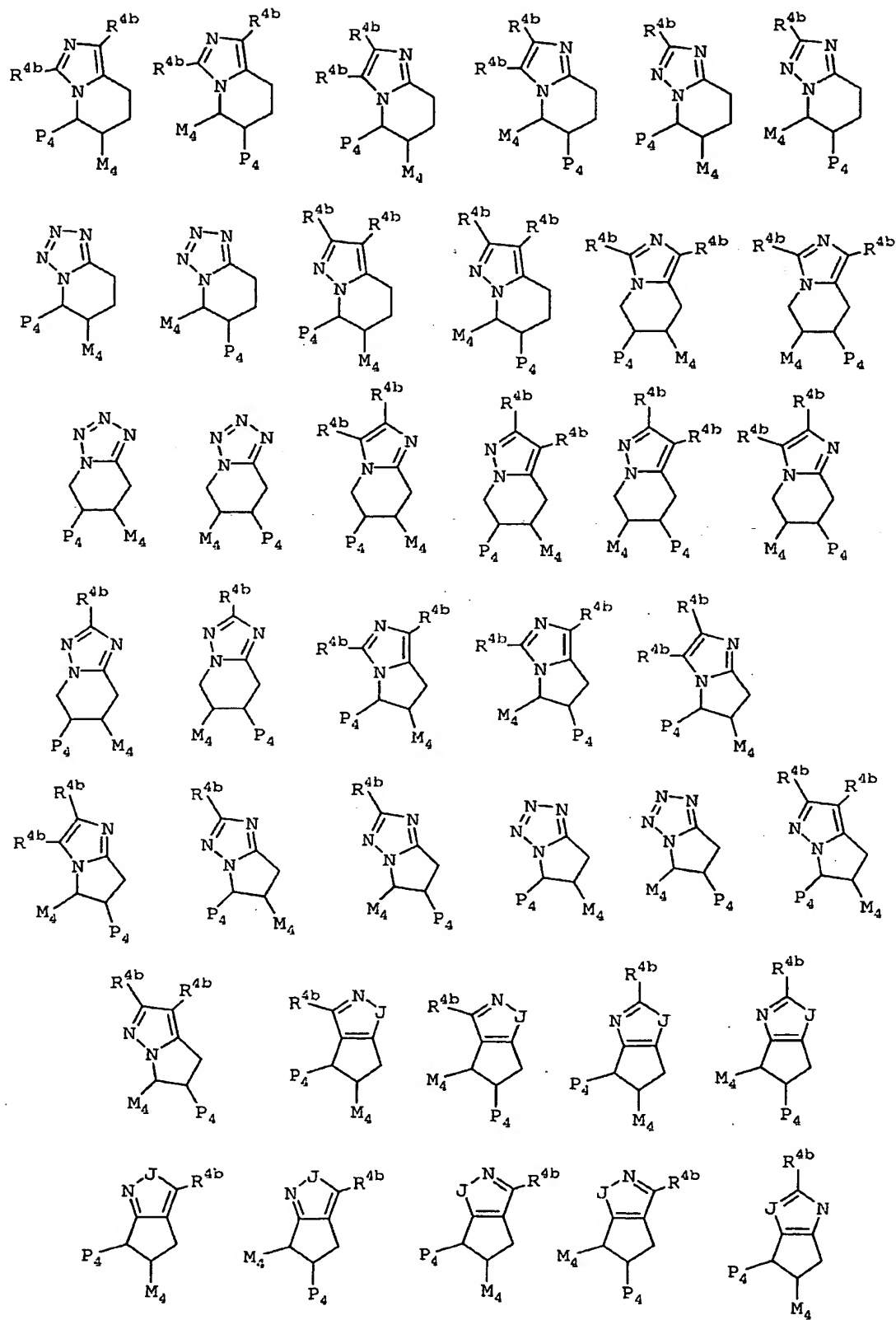


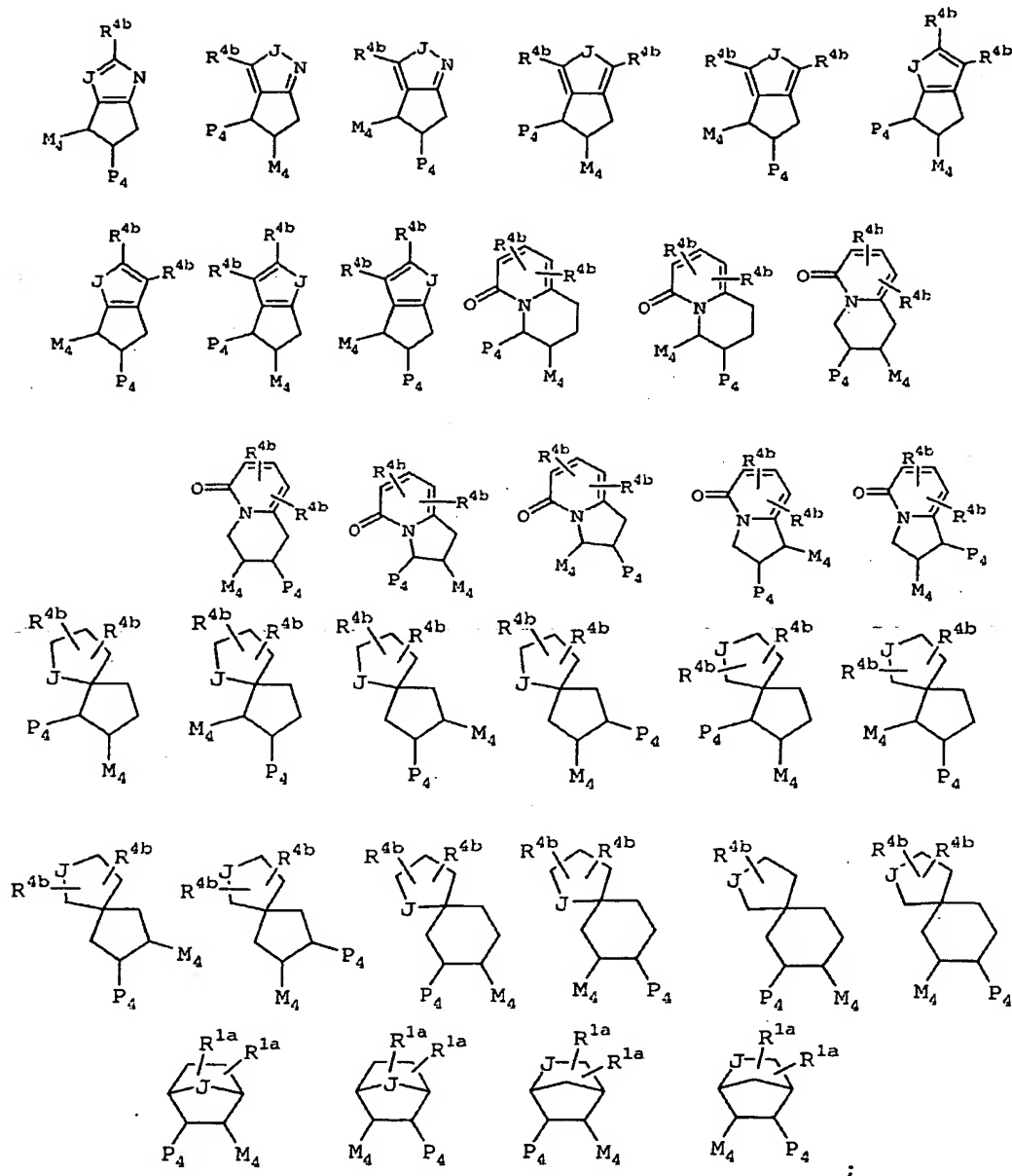










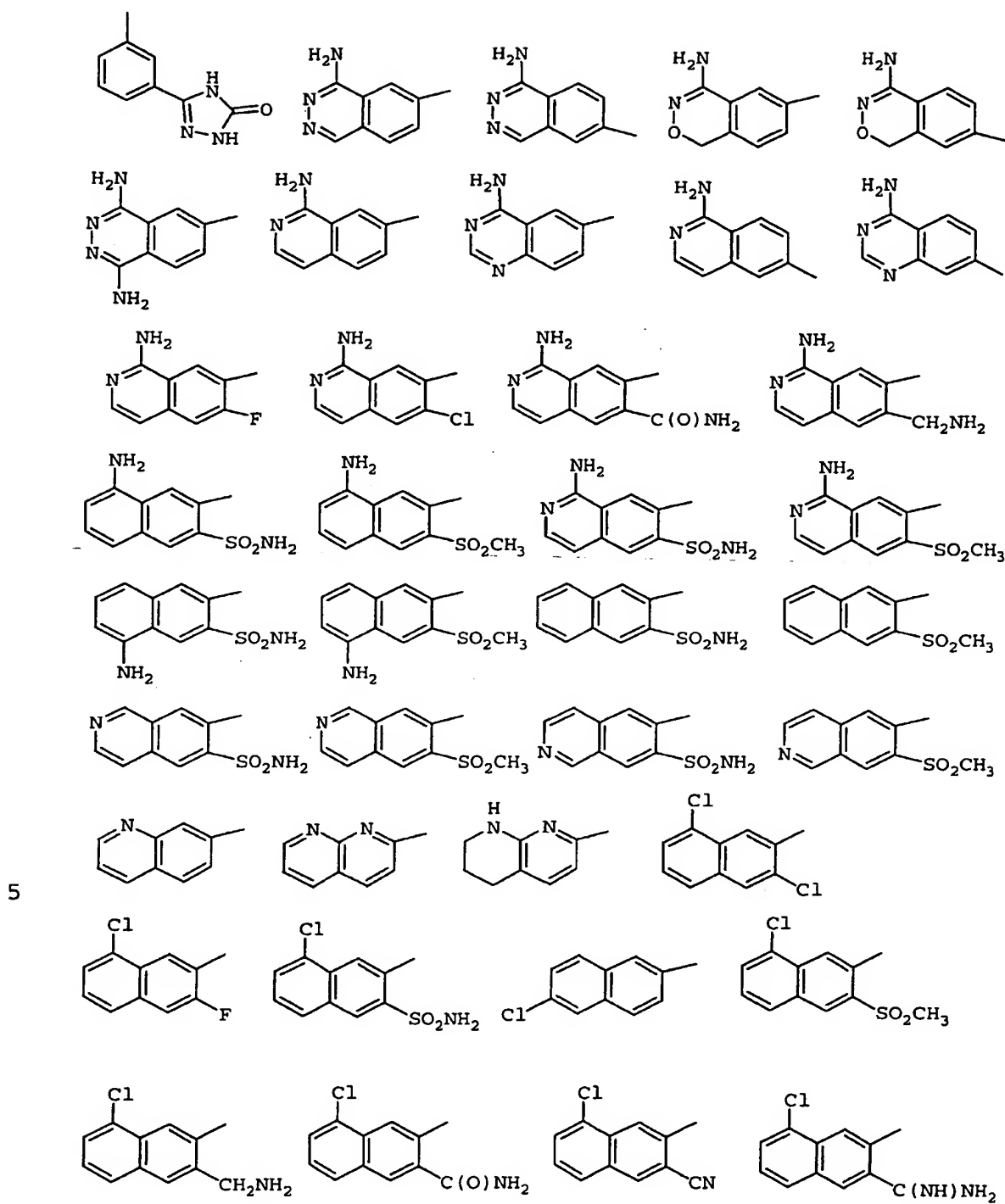


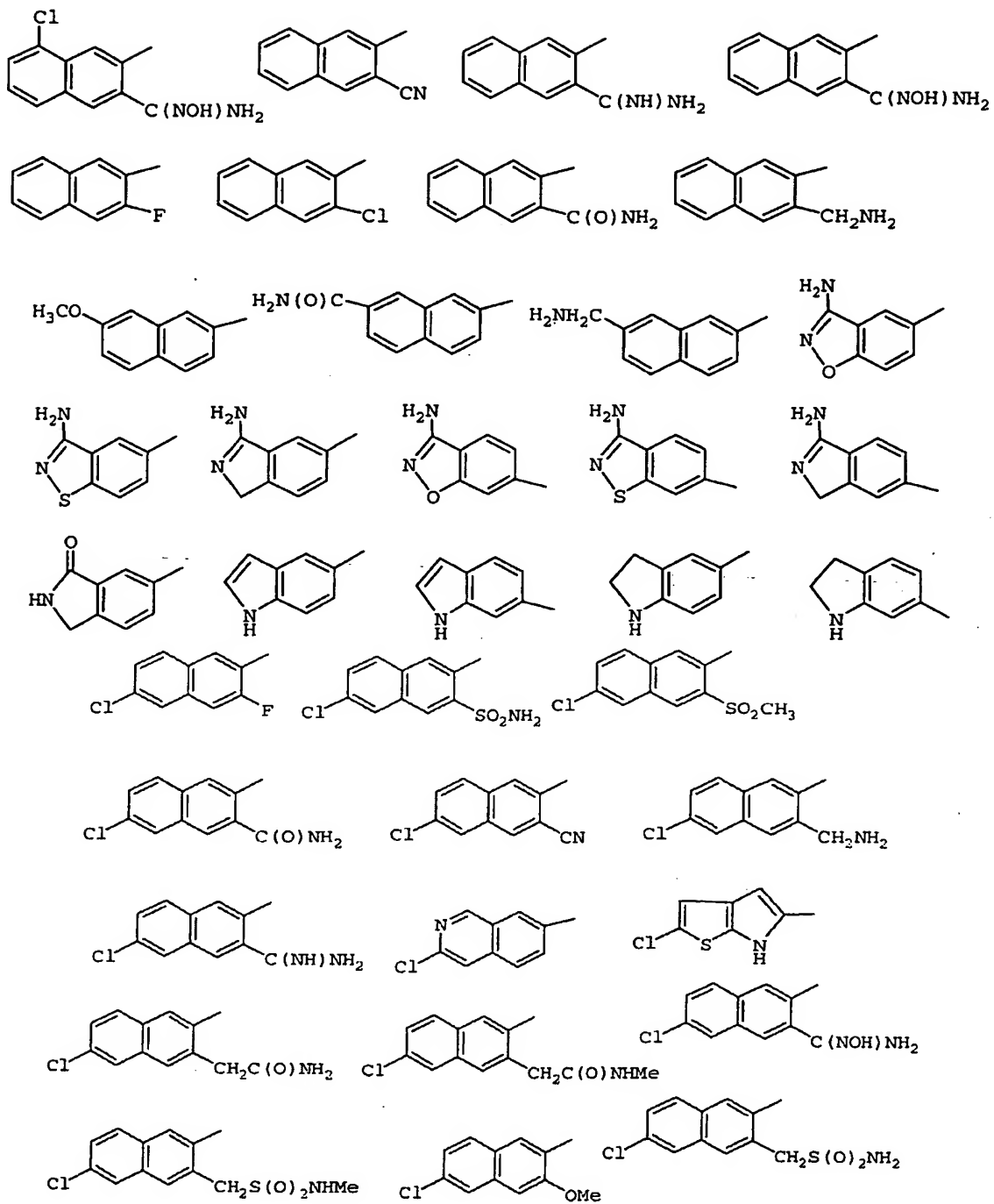
5 J is selected from O, S, NH, and NR^{1a};

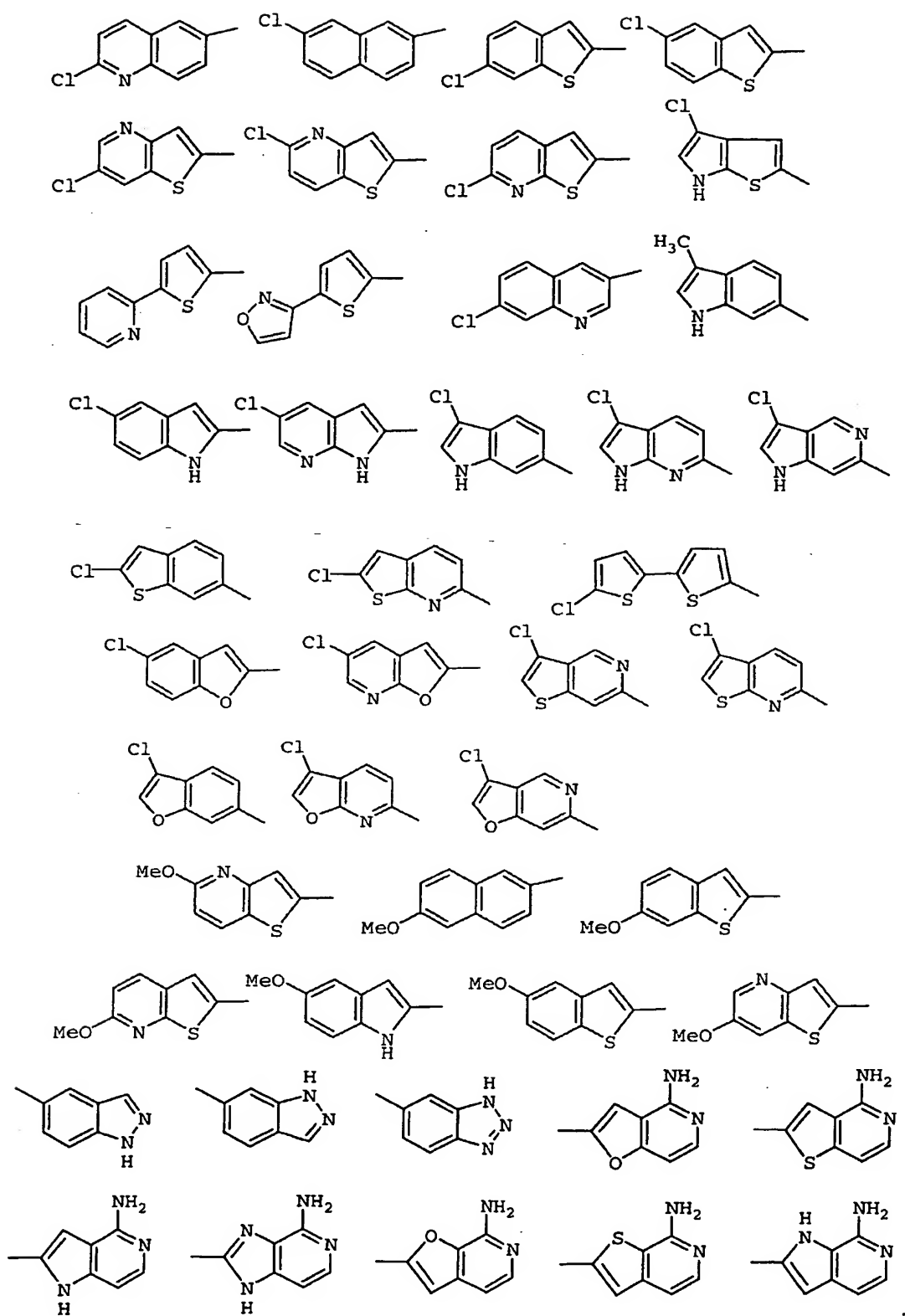
G is selected from the group:

- 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
- 2-aminomethyl-3-fluoro-phenyl;
- 10 2-aminomethyl-4-fluoro-phenyl;
- 2-aminomethyl-4-methoxy-phenyl;
- 2-aminomethyl-5-fluoro-phenyl;
- 2-aminomethyl-5-methoxy-phenyl;

- 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
2-aminosulfonyl-phenyl; 2-methylsulfonyl-phenyl;
3-(N,N-dimethylamino)-4-chloro-phenyl;
5 3-(N,N-dimethylamino)-phenyl;
3-(N-methylamino)-4-chloro-phenyl;
3-(N-methylamino)-phenyl; 3-amido-phenyl;
3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
3-amino-phenyl; 3-chloro-phenyl; 3,5-dichloro-thien-2-yl;
10 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
4-(N-methylamino)-5-chloro-thien-2-yl;
4-amino-5-chloro-thien-2-yl; 4-chloro-phenyl;
4-methoxy-2-methylsulfonyl-phenyl; 4-methoxy-phenyl;
5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
15 5-(N-methylamino)-4-chloro-thien-2-yl;
5-amino-4-chloro-thien-2-yl; 5-chloro-pyrid-2-yl;
5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-chloro-
pyrimidin-3-yl; 6-chloro-pyridazin-3-yl;
20 2-aminomethyl-4-chloro-phenyl;
2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl;
4-chloro-2-methylsulfonyl-phenyl;
2-aminosulfonyl-4-fluoro-phenyl; 2-amido-4-fluoro-phenyl;
4-fluoro-2-methylsulfonyl-phenyl;
25 2-aminomethyl-4-bromo-phenyl;
2-aminosulfonyl-4-bromo-phenyl; 2-amido-4-bromo-phenyl;
4-bromo-2-methylsulfonyl-phenyl;
2-aminomethyl-4-methyl-phenyl;
2-aminosulfonyl-4-methyl-phenyl; 2-amido-4-methyl-phenyl;
30 2-methylsulfonyl-4-methyl-phenyl; 4-fluoro-pyrid-2-yl;
4-bromo-pyrid-2-yl; 4-methyl-pyrid-2-yl;
5-fluoro-thien-2-yl; 5-bromo-thien-2-yl;
5-methyl-thien-2-yl; 2-amido-4-methoxy-phenyl;







G_1 is absent or is selected from CH_2 , CH_2CH_2 , $CH=CH$, CH_2O , OCH_2 , NH , CH_2NH , $NHCH_2$, $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $NHC(O)NH$, $C(O)NHS(O)_2$, $NHCOCONH$, $NHCOC(S)NH$, $NHC(S)CONH$, $CH_2S(O)_2$, $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$,
5 provided that G_1 does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;

A is selected from cyclohexyl, indoliny, piperidiny, phenyl, pyridyl, and pyrimidyl, and is substituted
10 with 0-2 R^4 ;

X is selected from CH_2 , $C(O)$, $-S(O)_2-$, $-NHC(O)-$, $-C(O)NH-$, $-CH_2NH-$, O , and $-CH_2O-$;

15 Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentanonyl, cyclohexyl, cyclohexanonyl, pyrrolidiny, pyrrolidinonyl, piperidiny, piperidinonyl, tetrahydrofuranyl, and tetrahydropyranyl, and, when Y is a ring, Y is
20 substituted with 0-1 R^4 ;

R^{1a} , at each occurrence, is selected from H, R^{1b} , $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, and CH_2R^{1b} , provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

25

R^{1b} is selected from CH_3 , CH_2CH_3 , F, Cl, Br, $-CN$, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , CO_2R^{2a} , $S(O)_pR^2$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle
30 consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

- R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-1 R^{4b} , benzyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- R^{2a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, benzyl, phenyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- R^{2b} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, C_{1-5} alkyl substituted with 0-3 R^{4b} , benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 4-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- R^{2c} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, phenyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of carbon atoms and from 1-4

heteroatoms selected from the group consisting of N, O, and S(O)_p;

5 R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

20 R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

30 R⁴, at each occurrence, is selected from OH, OR², CH₂OR², (CH₂)₂OR², F, Br, Cl, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, CF₃, and CF₂CF₃;

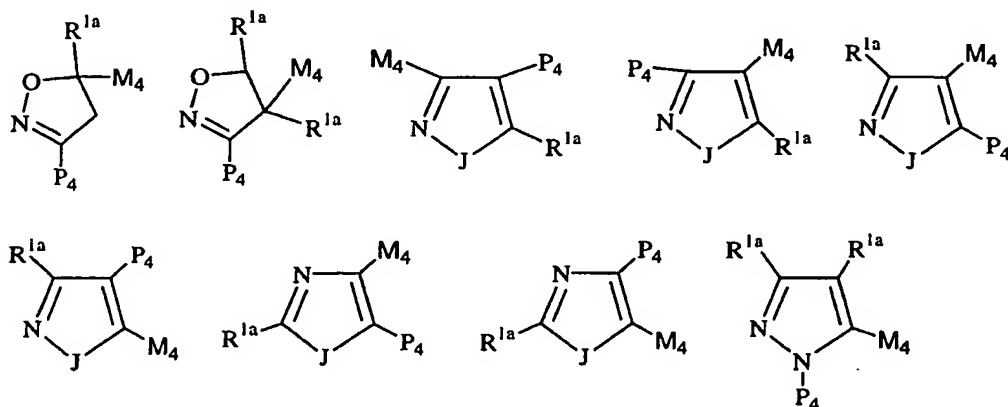
- R^{4a} is selected from $-(CR^3R^3g)_r$ -5-6 membered carbocycle substituted with 0-3 R^{4c} , $-(CR^3R^3g)_r$ -5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of:
- 5 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, $(CR^3R^3g)_rNR^{2d}R^{2d}$, $(CR^3R^3g)_rN(\rightarrow O)R^{2d}R^{2d}$, $(CR^3R^3g)_rOR^{2d}$, $(CR^3R^3g)_r-C(O)NR^{2d}R^{2d}$, $(CR^3R^3g)_r-NR^{2d}C(O)R^{2e}$, $(CR^3R^3g)_r-C(O)R^{2e}$, $(CR^3R^3g)_r-NR^{2d}C(O)NR^{2d}R^{2d}$, $(CR^3R^3g)_r-NR^{2d}C(O)OR^{2d}$, $(CR^3R^3g)_r-NR^{2d}SO_2R^{2d}$, and $(CR^3R^3g)_r-S(O)_pR^{2d}$, provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or $S(O)H$;
- R^{4b} , at each occurrence, is selected from H, $=O$, OR^3 ,
- 15 CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $-CN$, NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, NR^3SO_2 -phenyl, $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, and CF_3 ;
- R^{4c} , at each occurrence, is selected from $=O$, OR^2 , CH_2OR^2 , F, Br, Cl, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, C_{2-3} alkenyl, C_{2-3} alkynyl, $-CN$, NO_2 , $NR^{2d}R^{2a}$, $CH_2NR^{2d}R^{2a}$, $N(\rightarrow O)R^{2d}R^{2a}$, $CH_2N(\rightarrow O)R^{2d}R^{2a}$, $C(O)R^{2c}$, $CH_2C(O)R^{2c}$, $NR^{2d}C(O)R^{2b}$, $CH_2NR^{2d}C(O)R^{2b}$, $C(O)NR^{2d}R^{2a}$, $CH_2C(O)NR^{2d}R^{2a}$,
- 25 $SO_2NR^{2d}R^{2a}$, $CH_2SO_2NR^{2d}R^{2a}$, $NR^{2d}SO_2R^{5a}$, $CH_2NR^{2d}SO_2R^{5a}$, $S(O)_pR^{5a}$, $CH_2S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , C_{3-6} carbocycle substituted with 0-2 R^{4b} , $(CH_2)C_{3-6}$ carbocycle substituted with 0-2 R^{4b} , 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, and $(CH_2)5-6$ membered heterocycle substituted with 0-2 R^{4b} and consisting of
- 30

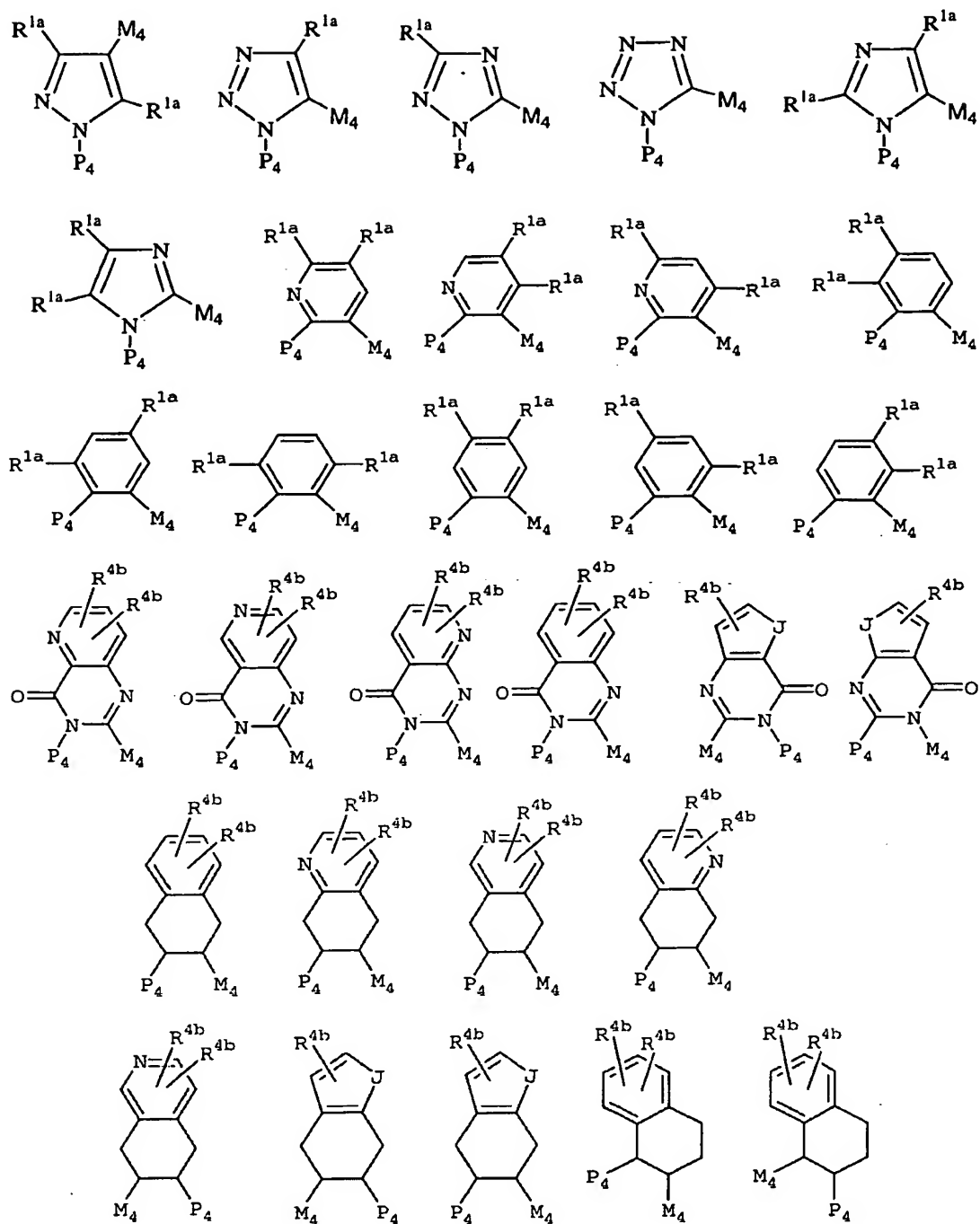
carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

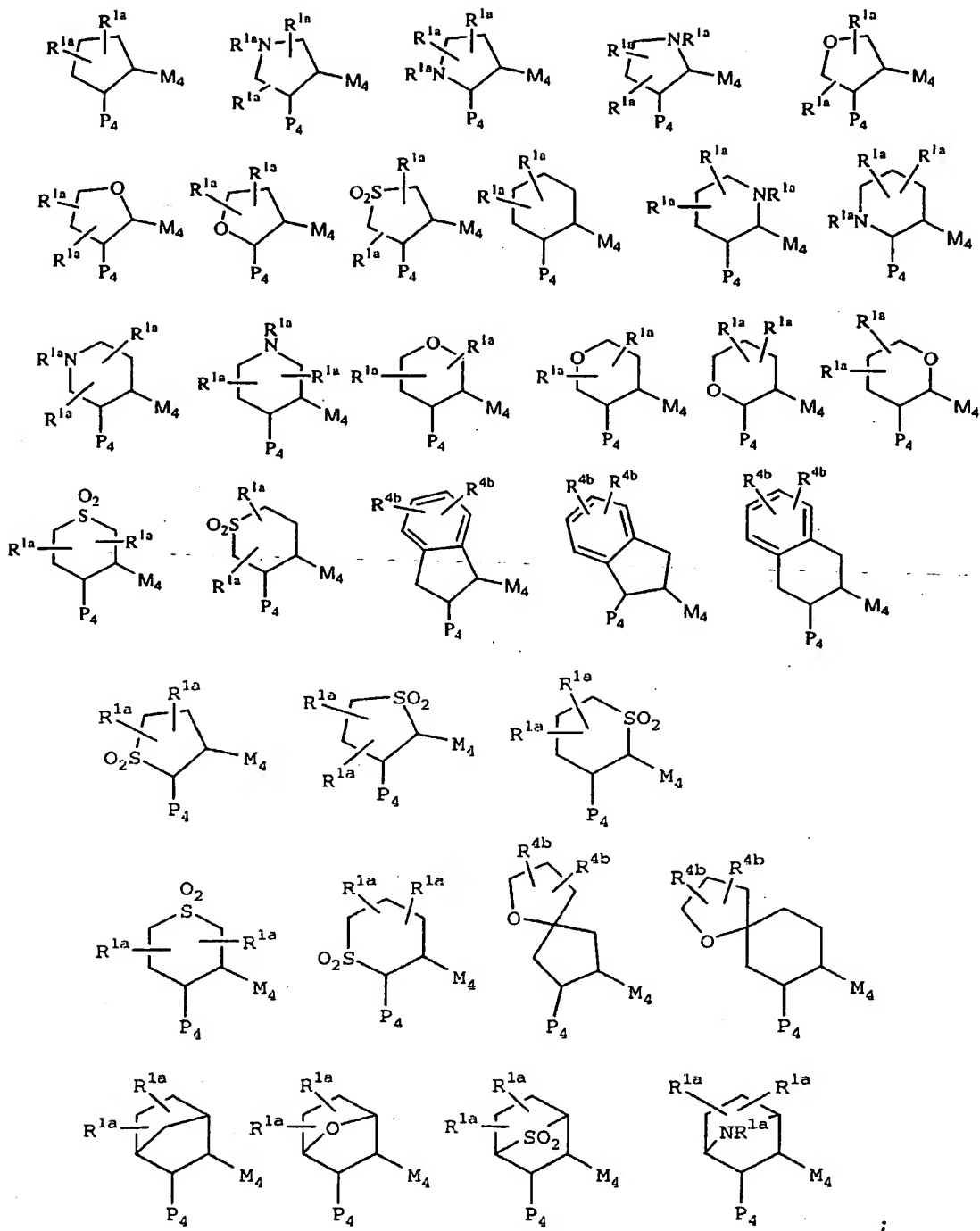
R⁵, at each occurrence, is selected from H, =O, CH₃, CH₂CH₃,
 5 CH₂CH₂CH₃, CH(CH₃)₂, OR³, CH₂OR³, F, Cl, -CN, NO₂,
 NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, C(O)OR^{3c}, NR³C(O)R^{3a},
 C(O)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, NR³SO₂-phenyl,
 S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl, CF₃, phenyl substituted
 with 0-2 R⁶, naphthyl substituted with 0-2 R⁶, and
 10 benzyl substituted with 0-2 R⁶; and,

R⁶, at each occurrence, is selected from H, OH, OR², F, Cl,
 CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, -CN, NO₂, NR²R^{2a},
 CH₂NR²R^{2a}, C(O)R^{2b}, CH₂C(O)R^{2b}, NR²C(O)R^{2b}, and
 15 SO₂NR²R^{2a}.

[12] In another preferred embodiment, the present invention
 provides a novel compound, wherein the compound is selected
 20 from:







J is selected from O, S, NH, and NR^{1a};

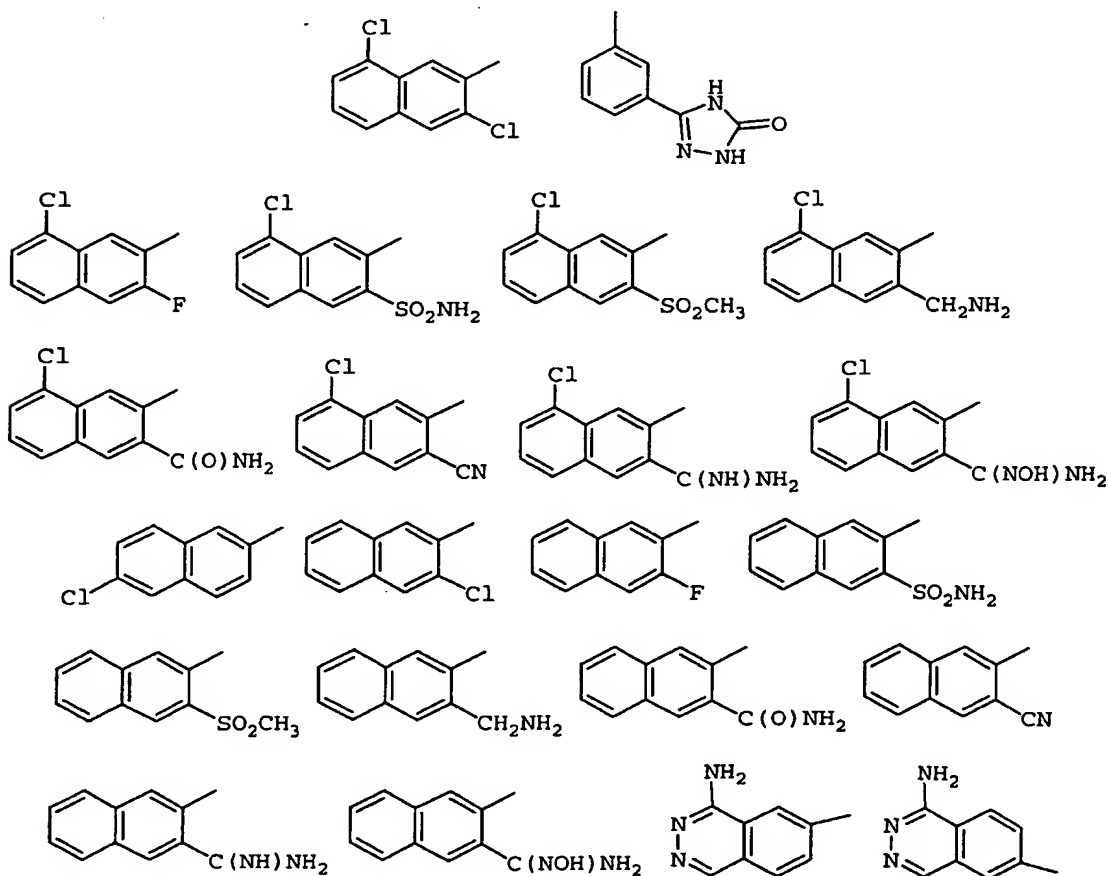
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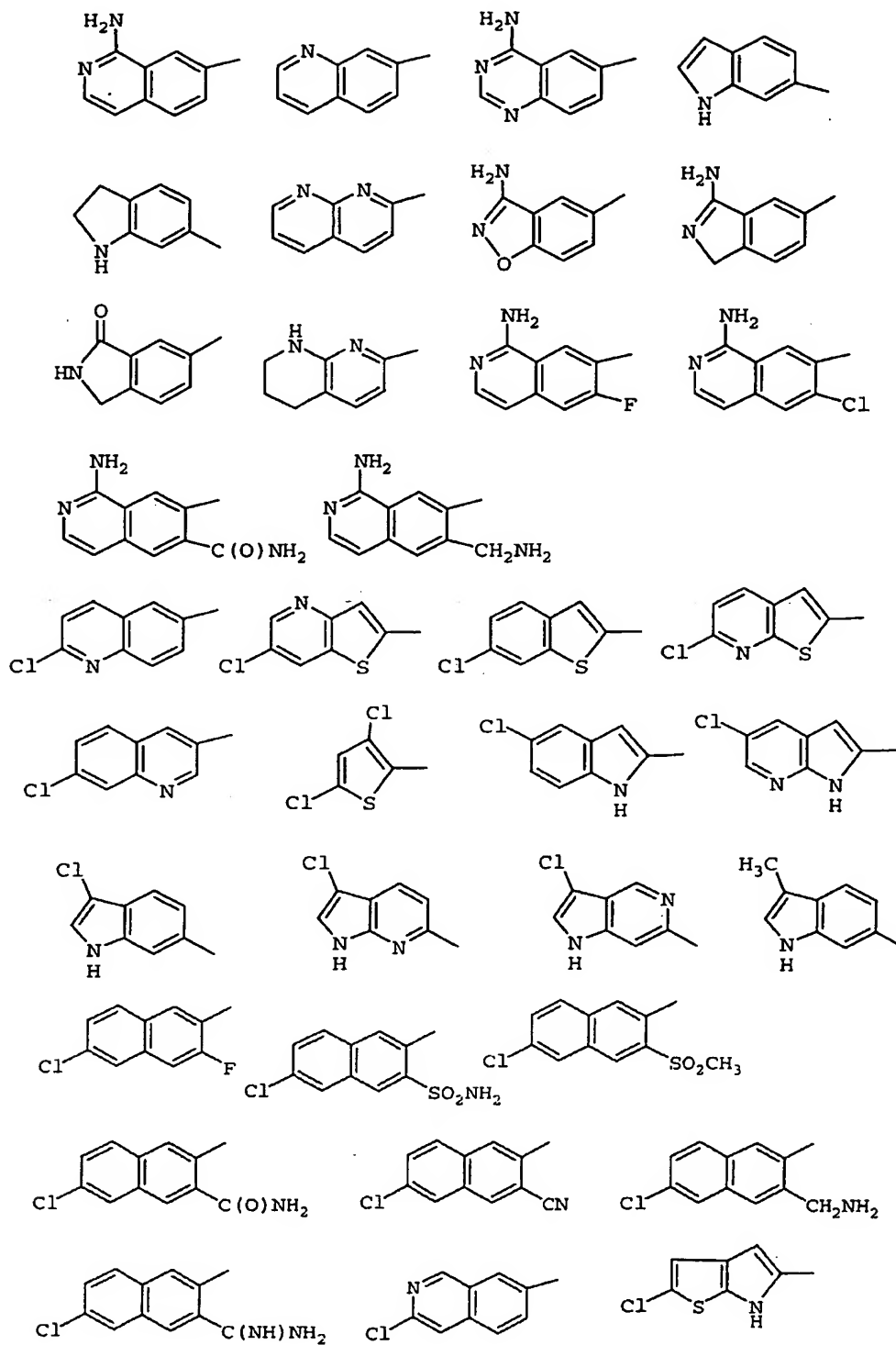
P₄ is -G₁-G;

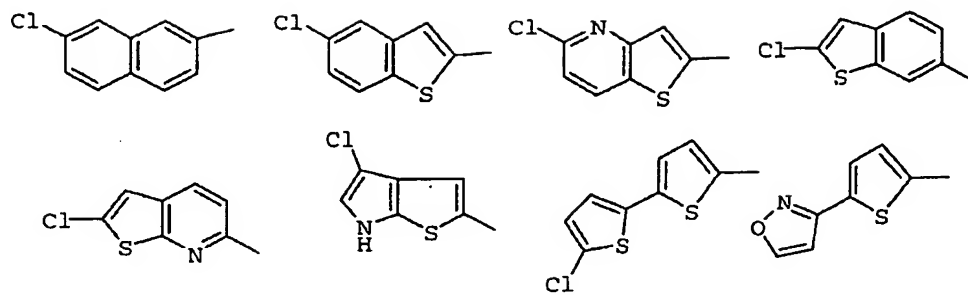
M₄ is -Z-A-B;

G is selected from:

- 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
 2-aminomethyl-3-fluoro-phenyl;
 5 2-aminomethyl-4-fluoro-phenyl;
 2-aminomethyl-5-fluoro-phenyl;
 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
 2-aminosulfonyl-phenyl; 3-amido-phenyl;
 10 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 3-chloro-phenyl; 4-chloro-phenyl; 4-methoxy-phenyl;
 5-chloro-pyrid-2-yl; 5-chloro-thien-2-yl;
 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-chloro-
 pyrimidin-3-yl; 6-chloro-pyridazin-3-yl;
 15 2-aminomethyl-4-chloro-phenyl;
 2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl;
 4-chloro-2-methylsulfonyl-phenyl;







G_1 is absent or is selected from $CH=CH$, CH_2NH , $NHCH_2$,
 $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $NHC(O)NH$, $CH_2S(O)_2$,
 5 $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$, provided that G_1 does not
 form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either
 group to which it is attached;

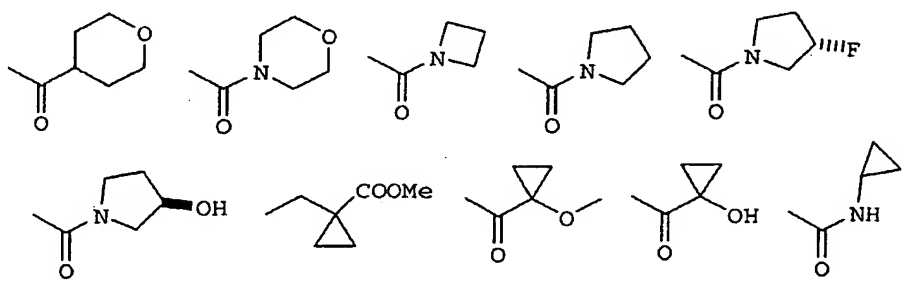
A is selected from the group: cyclohexyl, indolinyl,
 10 piperidinyl, phenyl, 2-pyridyl, 3-pyridyl, 2-
 pyrimidyl, 2-Cl-phenyl, 3-Cl-phenyl, 2-F-phenyl, 3-F-
 phenyl, 2-methylphenyl, 2-aminophenyl, and 2-
 methoxyphenyl;

15 Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl,
 cyclobutyl, cyclopentyl, 2-cyclopentanonyl,
 cyclohexyl, 2-cyclohexanonyl, pyrrolidinyl (attached
 to A and R^{4a} at the 2-position), pyrrolidinyl (attached
 to A and R^{4a} at the 3-position), 2-pyrrolidinonyl
 20 (attached to A and R^{4a} at the 3-position), piperidinyl
 (attached to A and R^{4a} at the 4-position), 4-
 piperidinonyl (attached to A and R^{4a} at the 3-position),
 tetrahydrofuranyl, and tetrahydropyranyl (attached to
 A and R^{4a} at the 4-position);

25 R^{1a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, CH_2F , CH_2Cl , Br, CH_2Br , $-CN$, CH_2CN , CF_3 ,
 CH_2CF_3 , OCH_3 , CH_2OH , $C(CH_3)_2OH$, CH_2OCH_3 , $CH_2CH_2OCH_3$,
 NH_2 , CH_2NH_2 , $NHCH_3$, CH_2NHCH_3 , $N(CH_3)_2$, $CH_2N(CH_3)_2$, CO_2H ,

$\text{CH}_2\text{CO}_2\text{H}$, $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, COCH_3 , CO_2CH_3 , $\text{CH}_2\text{CO}_2\text{CH}_3$, SCH_3 ,
 CH_2SCH_3 , $\text{S}(\text{O})\text{CH}_3$, $\text{CH}_2\text{S}(\text{O})\text{CH}_3$, $\text{S}(\text{O})_2\text{CH}_3$, $\text{CH}_2\text{S}(\text{O})_2\text{CH}_3$,
 $\text{C}(\text{O})\text{NH}_2$, $\text{CH}_2\text{C}(\text{O})\text{NH}_2$, SO_2NH_2 , $\text{CH}_2\text{SO}_2\text{NH}_2$, NHSO_2CH_3 ,
 $\text{CH}_2\text{NHSO}_2\text{CH}_3$, $\text{COCH}_2\text{C}(\text{CH}_3)_3$, COCH_2OH , $\text{COCH}_2\text{OCH}_3$,
5 $\text{COC}(\text{CH}_3)_2\text{OH}$, $\text{COC}(\text{CH}_3)_2\text{CH}_2\text{OH}$, $\text{COC}(\text{CH}_3)_2\text{CH}_2\text{OCH}_3$,
 $\text{C}(\text{O})\text{OCH}_2\text{CH}_2\text{OCH}_3$, COCF_3 , $\text{CO}_2\text{CH}_2\text{CH}_3$, $\text{CO}_2\text{CH}(\text{CH}_3)_2$,
 $\text{CO}_2\text{C}(\text{CH}_3)_3$, $\text{CH}_2\text{CH}_2\text{CO}_2\text{CH}_2\text{CH}_3$, $\text{CONH}(\text{CH}_3)$, $\text{CONH}(\text{CH}_2\text{CH}_3)$,
 $\text{CONHC}(\text{CH}_3)_3$, $\text{CON}(\text{CH}_3)_2$, $\text{CON}(\text{CH}_3)(\text{CH}_2\text{CH}_3)$,
 $\text{CON}(\text{CH}_3)\text{CH}(\text{CH}_3)_2$, $\text{CH}_2\text{CON}(\text{CH}_3)_2$, $\text{C}(\text{O})$ -phenyl, $\text{C}(\text{O})$ -
10 cyclopropyl, $\text{C}(\text{O})$ -cyclobutyl, $\text{C}(\text{O})$ -cyclopentyl,
pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyridin-2-
yl-N-oxide, pyridin-3-yl-N-oxide, pyridin-4-yl-N-
oxide, imidazol-1-yl, CH_2 -imidazol-1-yl, 4-methyl-
oxazol-2-yl, 4-N,N-dimethylaminomethyl-oxazol-2-yl,
15 1,2,3,4-tetrazol-1-yl, 1,2,3,4-tetrazol-5-yl, CH_2 -
1,2,3,4-tetrazol-1-yl, and CH_2 -1,2,3,4-tetrazol-5-yl,
provided that R^{1a} forms other than an N-halo, N-S, or
N-CN bond;

20 alternatively, R^{1a} is selected from:



R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 ,
 $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, phenyl substituted with 0-1 R^{4b} ,
25 benzyl substituted with 0-1 R^{4b} , and 5 membered
aromatic heterocycle substituted with 0-1 R^{4b} and
consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and $\text{S}(\text{O})_p$;

R^{2a}, at each occurrence, is selected from H, CH₃, and CH₂CH₃;

5 alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group
10 consisting of N, O, and S(O)_p;

R^{2b}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, CH₃, and CH₂CH₃;

15 R^{2c}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, CH₃, and CH₂CH₃;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl substituted with 0-2 R^{4c}, and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c} consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;
20
25

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl substituted with 0-2 R^{4c}, and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p,
30

provided that R^{2e} forms other than a $C(O)$ -halo or $C(O)$ - $S(O)_p$ moiety;

- R^{4a} is selected from $-(CH_2)_r$ -5-6 membered carbocycle substituted with 0-3 R^{4c} , $-(CH_2)_r$ -5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, $(CH_2)_rNR^{2d}R^{2d}$, $(CH_2)_rN(\rightarrow O)R^{2d}R^{2d}$, $(CH_2)_rOR^{2d}$, $(CH_2)_rC(O)NR^{2d}R^{2d}$, $(CH_2)_rNR^{2d}C(O)R^{2e}$, $(CH_2)_rC(O)R^{2e}$, $(CH_2)_rNR^{2d}C(O)NR^{2d}R^{2d}$, $(CH_2)_rNR^{2d}C(O)OR^{2d}$, $(CH_2)_rNR^{2d}SO_2R^{2d}$, and $(CH_2)_rS(O)_pR^{2d}$, provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or $S(O)H$;
- R^{4b} , at each occurrence, is selected from H, $=O$, OR^3 , CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, NR^3SO_2 -phenyl, $S(O)_2CH_3$, $S(O)_2$ -phenyl, and CF_3 ;
- R^{4c} , at each occurrence, is selected from $=O$, OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, C_{2-3} alkenyl, C_{2-3} alkynyl, CH_2OH , CH_2OCH_3 , $CH_2OCH_2CH_3$, $CH_2OCH_2CH_2CH_3$, $CH_2OCH(CH_3)_2$, F, Br, Cl, CF_3 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $N(\rightarrow O)R^2R^{2a}$, $CH_2N(\rightarrow O)R^2R^{2a}$, $C(O)R^{2c}$, $CH_2C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $CH_2NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $CH_2C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $CH_2SO_2NR^2R^{2a}$, $NR^2SO_2R^{5a}$, $CH_2NR^2SO_2R^{5a}$, $S(O)_pR^{5a}$, $CH_2S(O)_pR^{5a}$, CF_3 , cyclopropyl substituted with 0-1 R^{4b} , cyclobutyl substituted with 0-1 R^{4b} , cyclopentyl substituted with 0-1 R^{4b} , phenyl substituted with 0-1 R^{4b} , $-CH_2$ -cyclopropyl substituted with 0-1 R^{4b} , $-CH_2$ -cyclobutyl substituted with 0-1 R^{4b} ,

-CH₂-cyclopentyl substituted with 0-1 R^{4b}, benzyl substituted with 0-2 R^{4b}, 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, and (CH₂)₅₋₆ membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

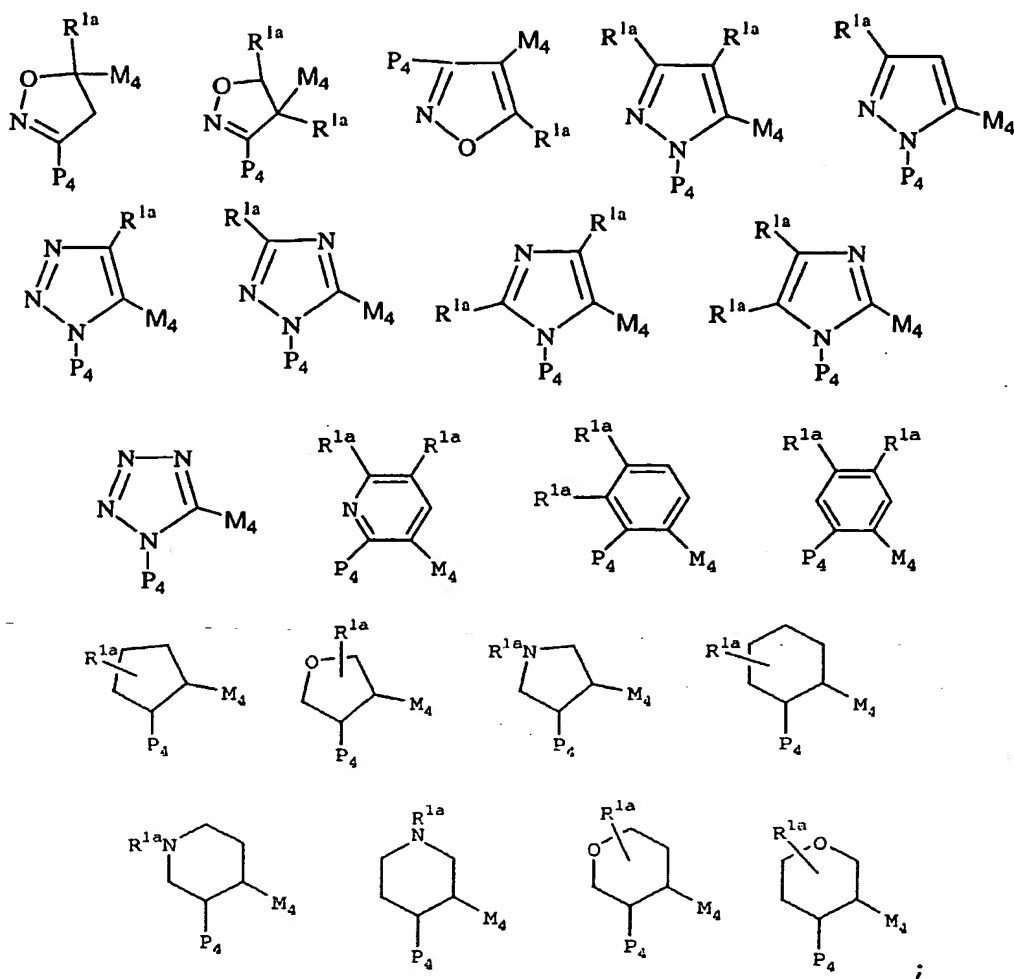
10 R⁵, at each occurrence, is selected from H, =O, CH₃, CH₂CH₃, OR³, CH₂OR³, F, Cl, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, C(O)OR^{3c}, NR³C(O)R^{3a}, C(O)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, NR³SO₂-phenyl, S(O)₂-CH₃, S(O)₂-phenyl, CF₃, phenyl

15 substituted with 0-2 R⁶, naphthyl substituted with 0-2 R⁶, and benzyl substituted with 0-2 R⁶; and,

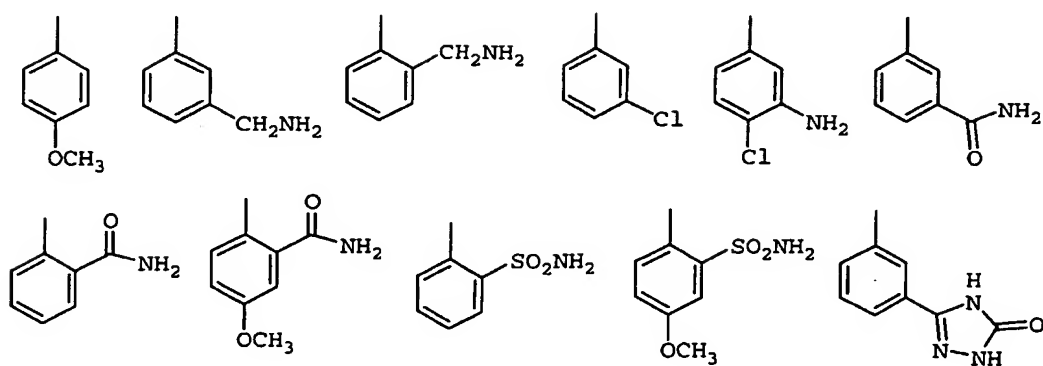
20 R⁶, at each occurrence, is selected from H, OH, OR², F, Cl, CH₃, CH₂CH₃, NR²R^{2a}, CH₂NR²R^{2a}, C(O)R^{2b}, CH₂C(O)R^{2b}, NR²C(O)R^{2b}, and SO₂NR²R^{2a}.

[13] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is selected from:

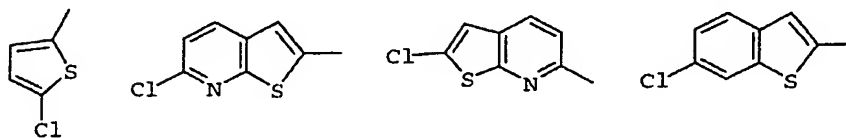
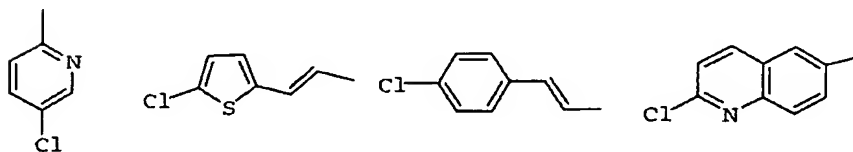
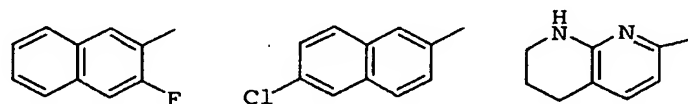
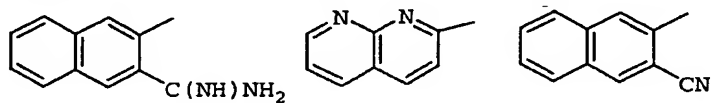
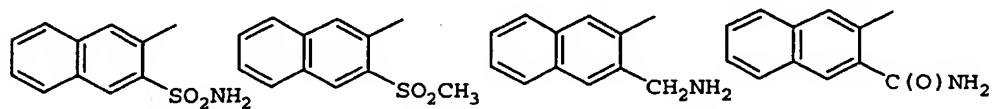
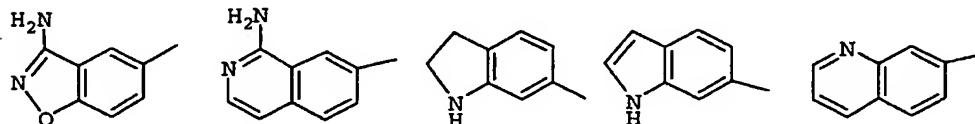
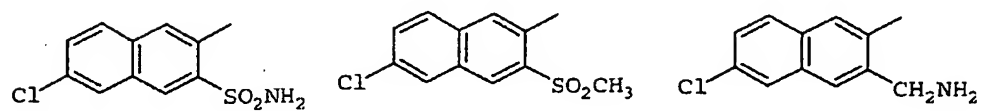
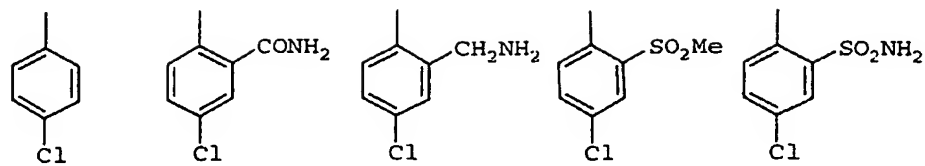
25

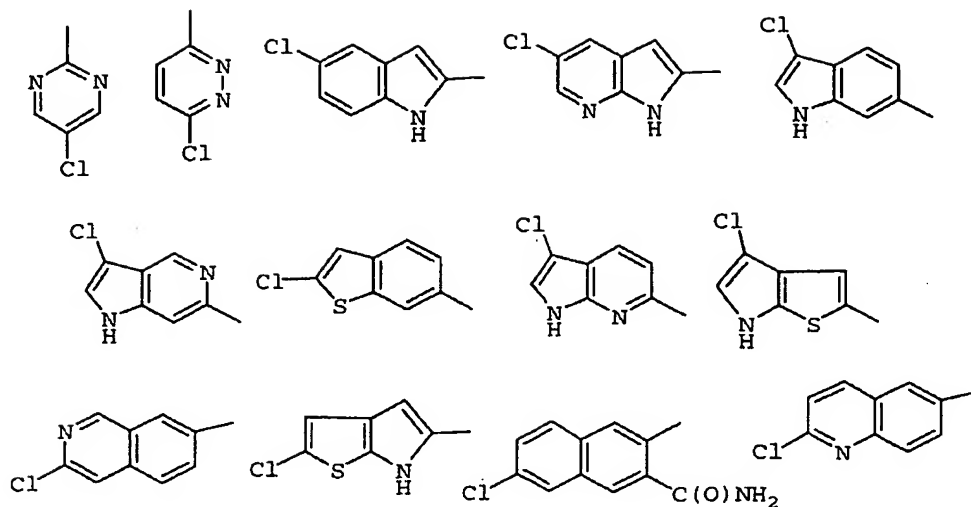


-G₁-G is selected from:

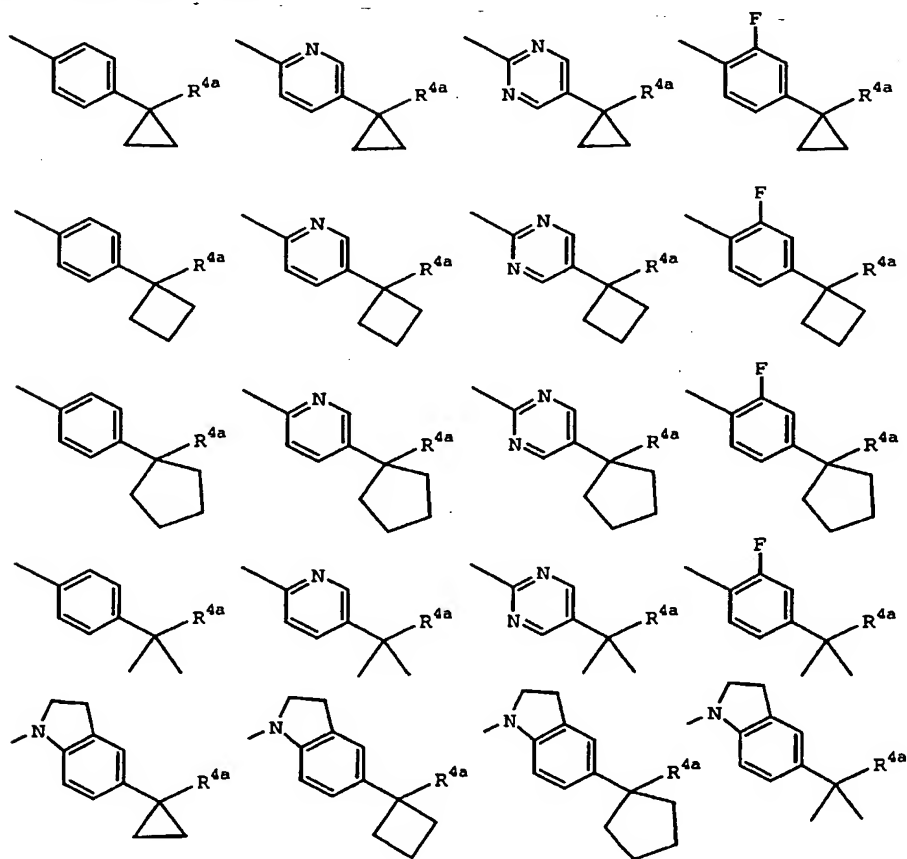


5





A-B is selected from:



5

R^{2d}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted

with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and a
 5-6 membered aromatic heterocycle consisting of:
 carbon atoms and 1-4 heteroatoms selected from the
 group consisting of N, O, and $S(O)_p$, provided that R^{2d}
 5 forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-
 halo, N-S, S-N, $S(O)_p$ - $S(O)_p$, S-O, O-N, O-S, or O-O
 moiety;

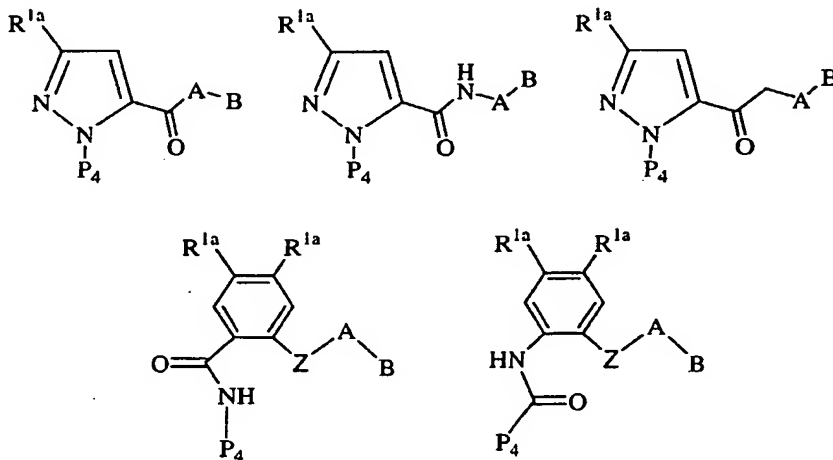
R^{2e} , at each occurrence, is selected from H, C_{1-4} alkyl
 10 substituted with 0-1 R^{4c} , C_{3-6} cycloalkyl substituted
 with 0-2 R^{4c} , phenyl, substituted with 0-2 R^{4c} , and 5-6
 membered aromatic heterocycle consisting of: carbon
 atoms and 1-4 heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$, provided that R^{2e} forms
 15 other than a $C(O)$ -halo or $C(O)$ - $S(O)_p$ moiety;

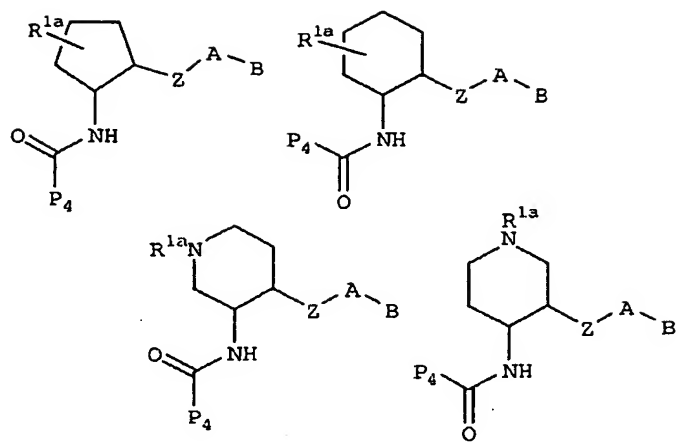
R^{4a} is selected from $NR^{2d}R^{2d}$, $CH_2NR^{2d}R^{2d}$, $CH_2CH_2NR^{2d}R^{2d}$,
 $N(\rightarrow O)R^{2d}R^{2d}$, $CH_2N(\rightarrow O)R^{2d}R^{2d}$, CH_2OR^{2d} , $C(O)R^{2e}$,
 $C(O)NR^{2d}R^{2d}$, $CH_2C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)R^{2e}$, $CH_2NR^{2d}C(O)R^{2e}$,
 20 $NR^{2d}C(O)NR^{2d}R^{2d}$, $CH_2NR^{2d}C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)OR^{2d}$,
 $CH_2NR^{2d}C(O)OR^{2d}$, $NR^{2d}SO_2R^{2d}$, $CH_2NR^{2d}SO_2R^{2d}$, $S(O)_pR^{2d}$,
 $CH_2S(O)_pR^{2d}$, 5-6 membered carbocycle substituted with
 0-2 R^{4c} , $-(CH_2)$ -5-6 membered carbocycle substituted
 with 0-2 R^{4c} , $-(CH_2)_2$ -5-6 membered carbocycle
 25 substituted with 0-2 R^{4c} , 5-6 membered heterocycle
 substituted with 0-2 R^{4c} and consisting of: carbon
 atoms and 1-4 heteroatoms selected from the group
 consisting of N, O, and $S(O)_p$, $-(CH_2)$ -5-6 membered
 heterocycle substituted with 0-2 R^{4c} and consisting of:
 30 carbon atoms and 1-4 heteroatoms selected from the
 group consisting of N, O, and $S(O)_p$, and $-(CH_2)_2$ -5-6
 membered heterocycle substituted with 0-2 R^{4c} and
 consisting of: carbon atoms and 1-4 heteroatoms

selected from the group consisting of N, O, and S(O)_p provided that S(O)_pR^{2d} forms other than S(O)₂H or S(O)H; and,

- 5 R^{4c} is selected from =O, OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH=CH₂, CH≡CH, CH₂OH, CH₂OCH₃, CH₂OCH₂CH₃, CH₂OCH₂CH₂CH₃, CH₂OCH(CH₃)₂, F, Br, Cl, CF₃, NR²R^{2a}, CH₂NR²R^{2a}, C(O)R^{2c}, CH₂C(O)R^{2c}, NR²C(O)R^{2b}, CH₂NR²C(O)R^{2b},
 10 C(O)NR²R^{2a}, CH₂C(O)NR²R^{2a}, SO₂NR²R^{2a}, CH₂SO₂NR²R^{2a}, NR²SO₂R^{5a}, CH₂NR²SO₂R^{5a}, S(O)_pR^{5a}, and CH₂S(O)_pR^{5a}.

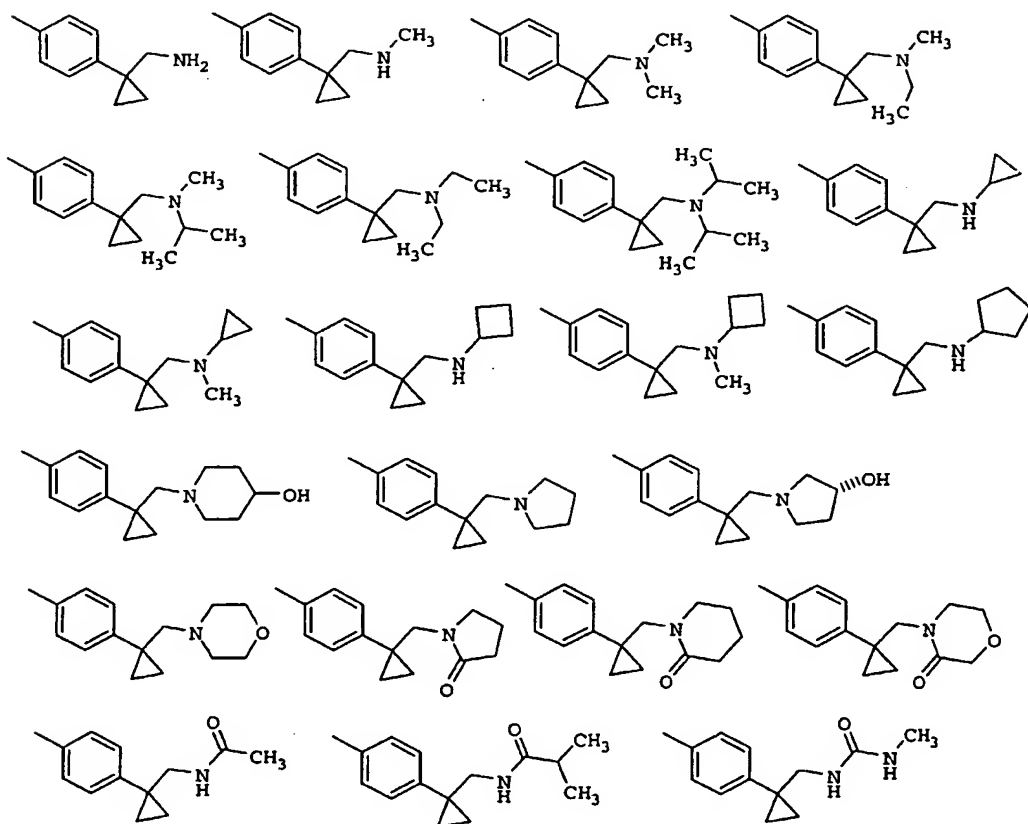
[14] In another preferred embodiment, the present invention
 15 provides a novel compound, wherein the compound is selected from:

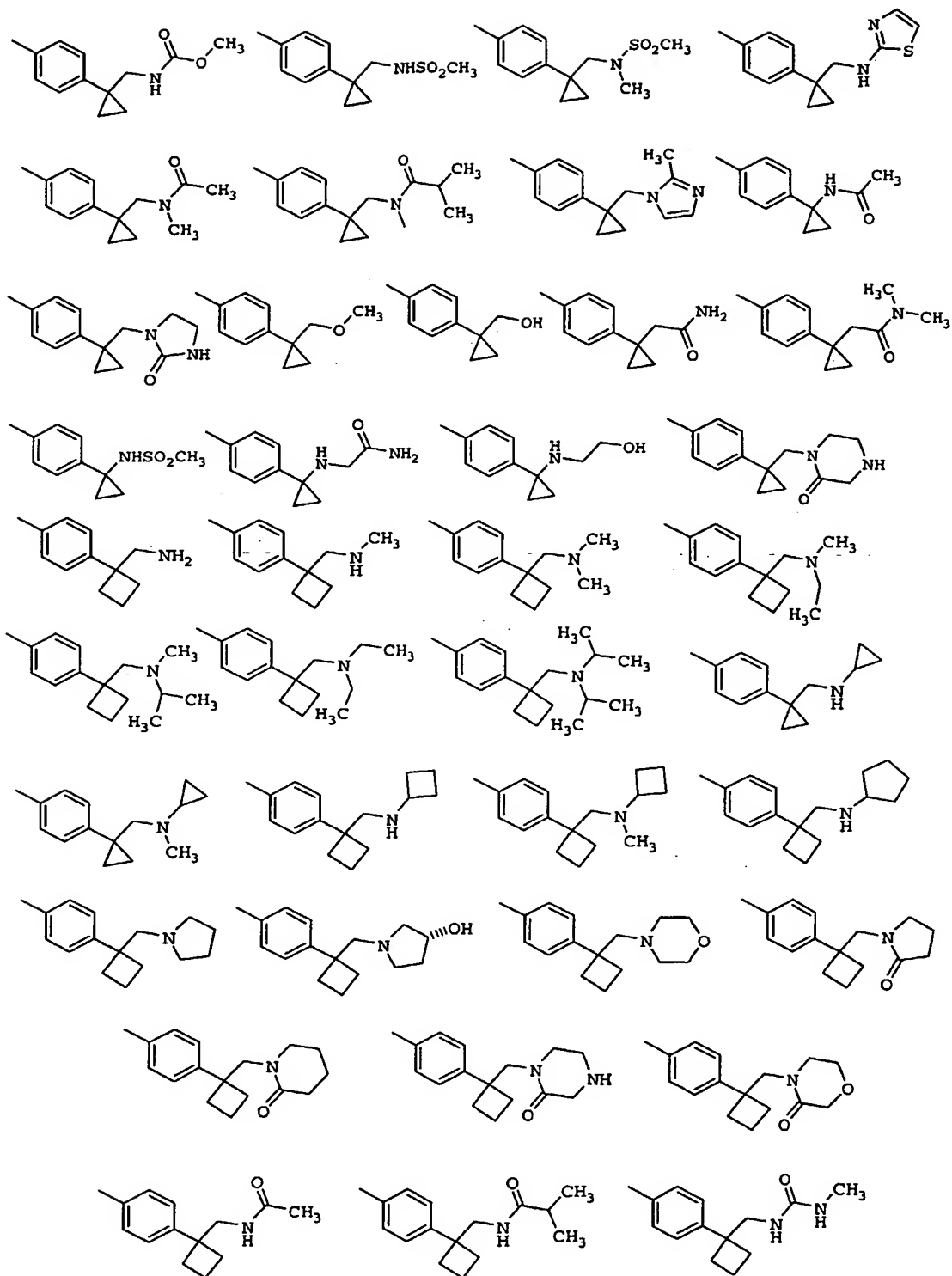


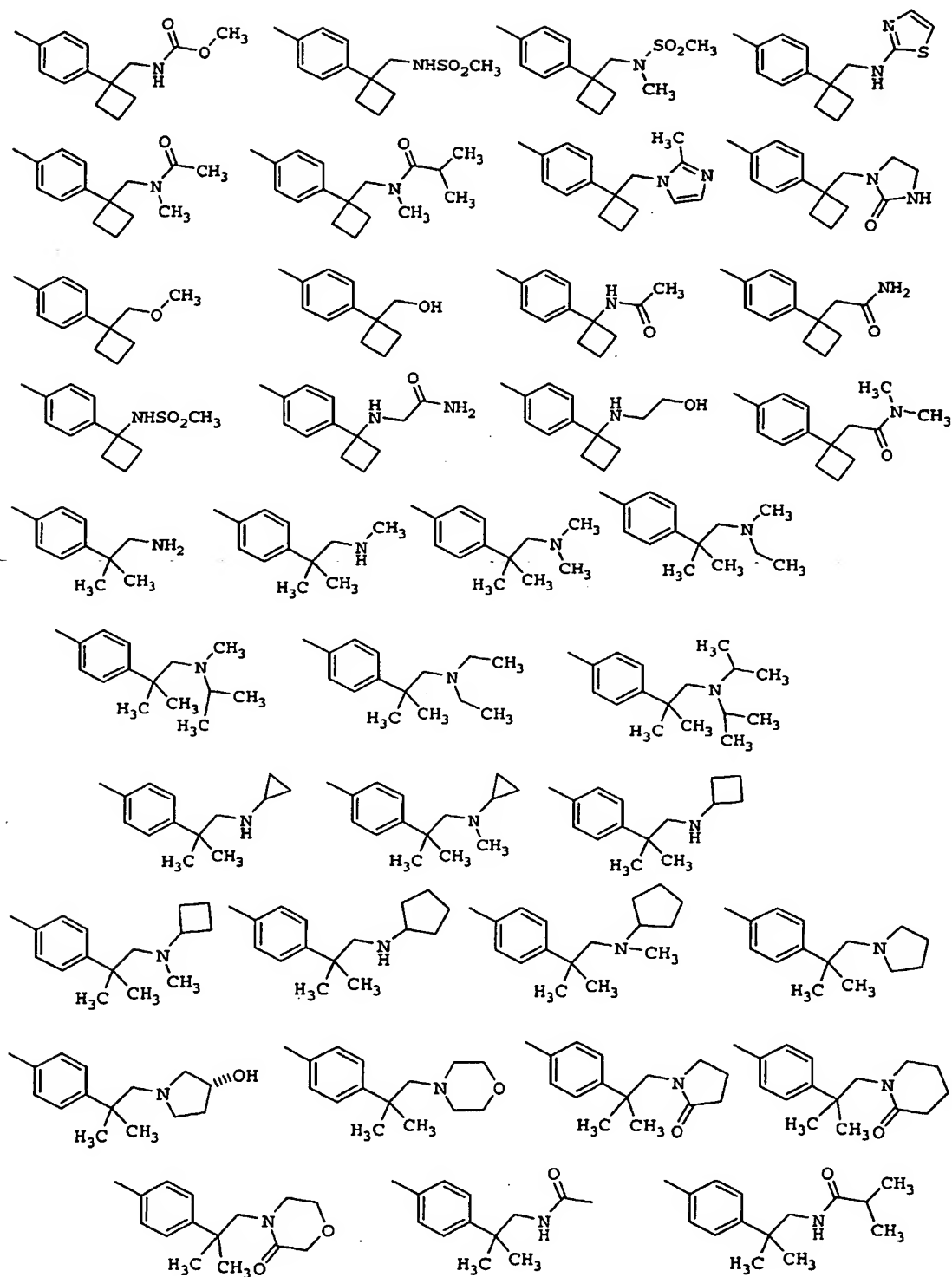


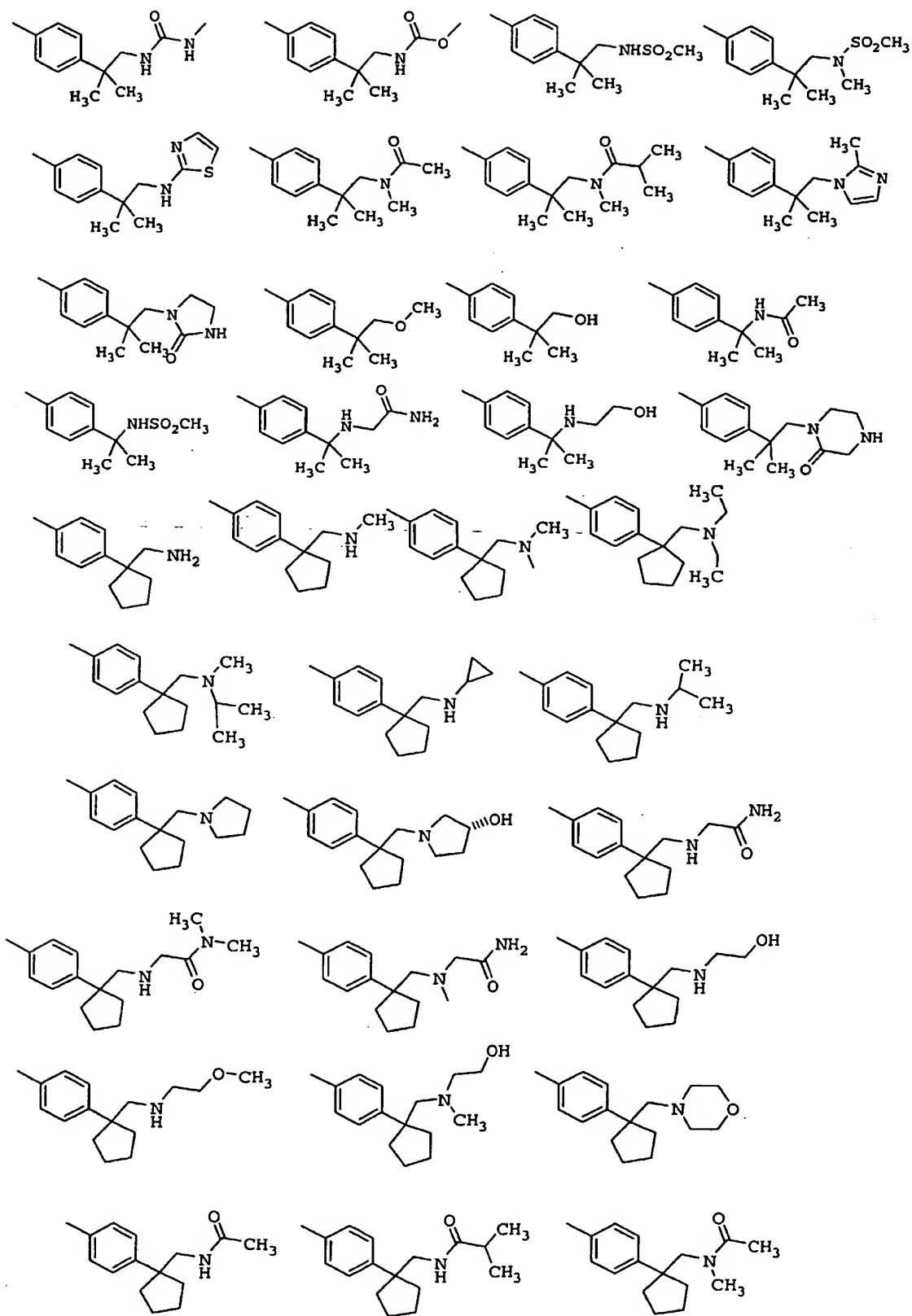
Z is selected from a $NHCH_2$, $C(O)NH$, $NHC(O)$, and $NHSO_2$; and,

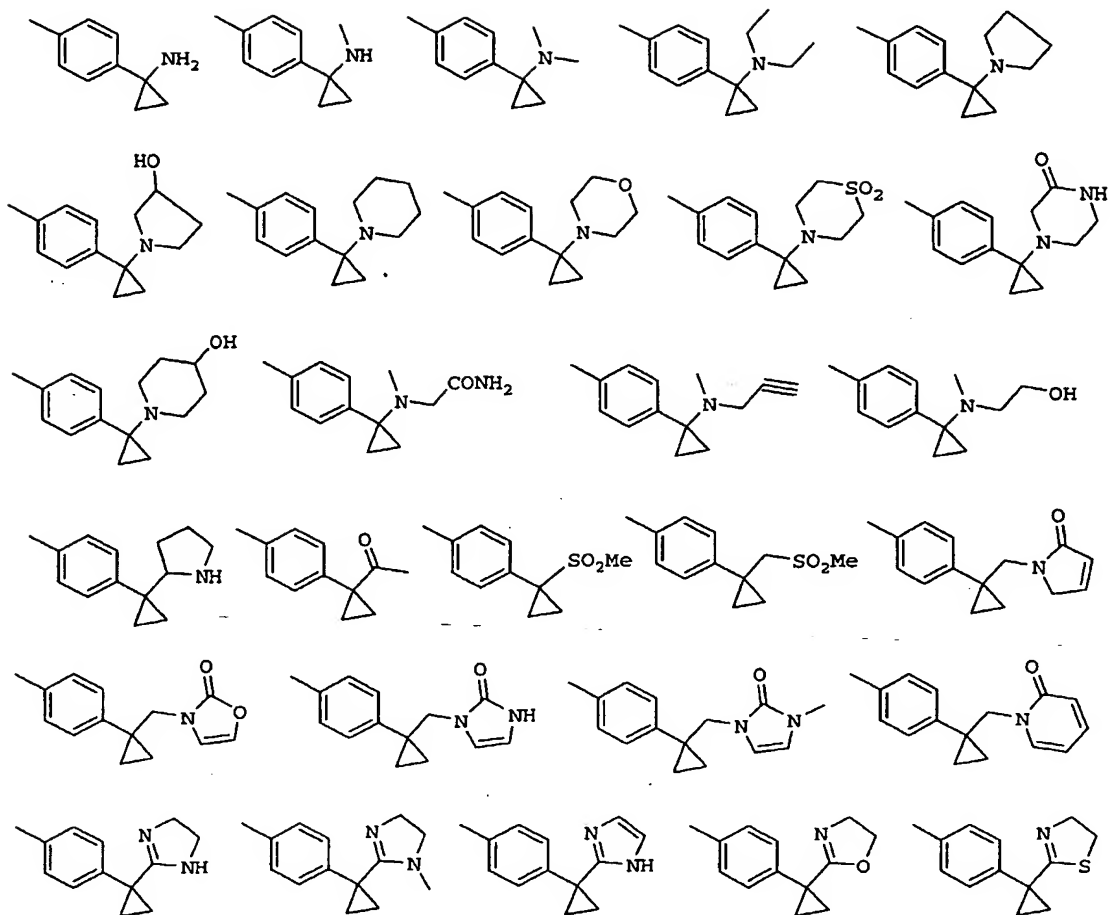
5 $A-B$ is selected from:

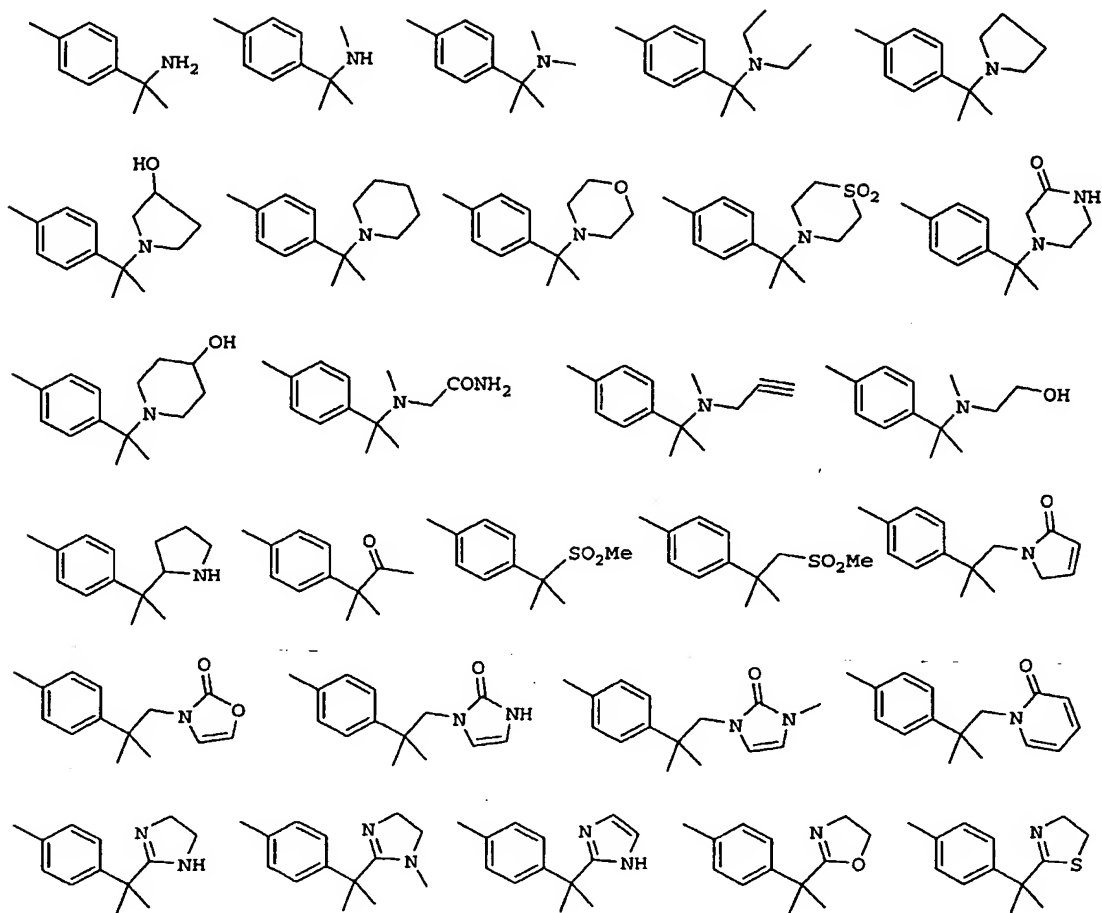


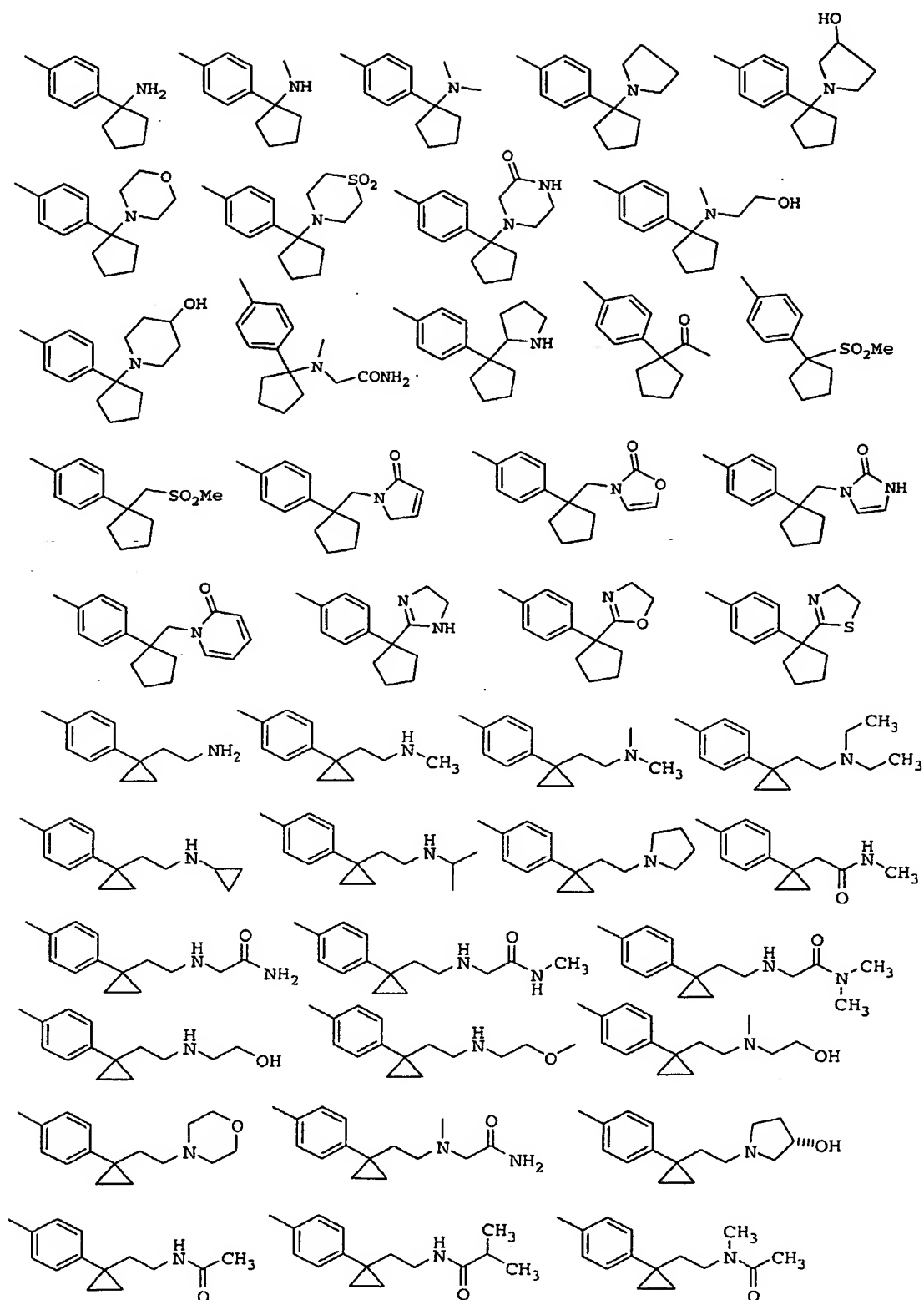


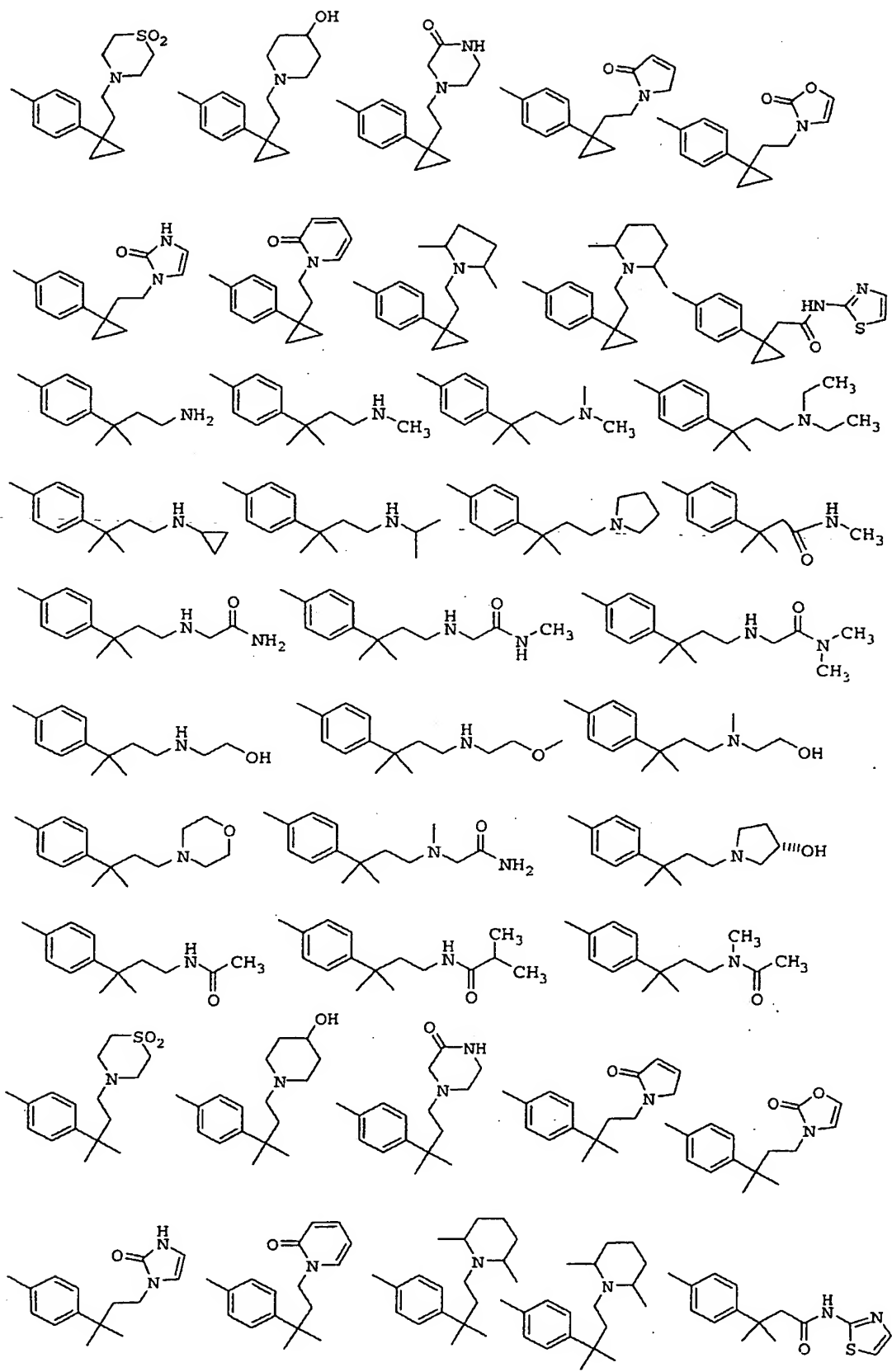


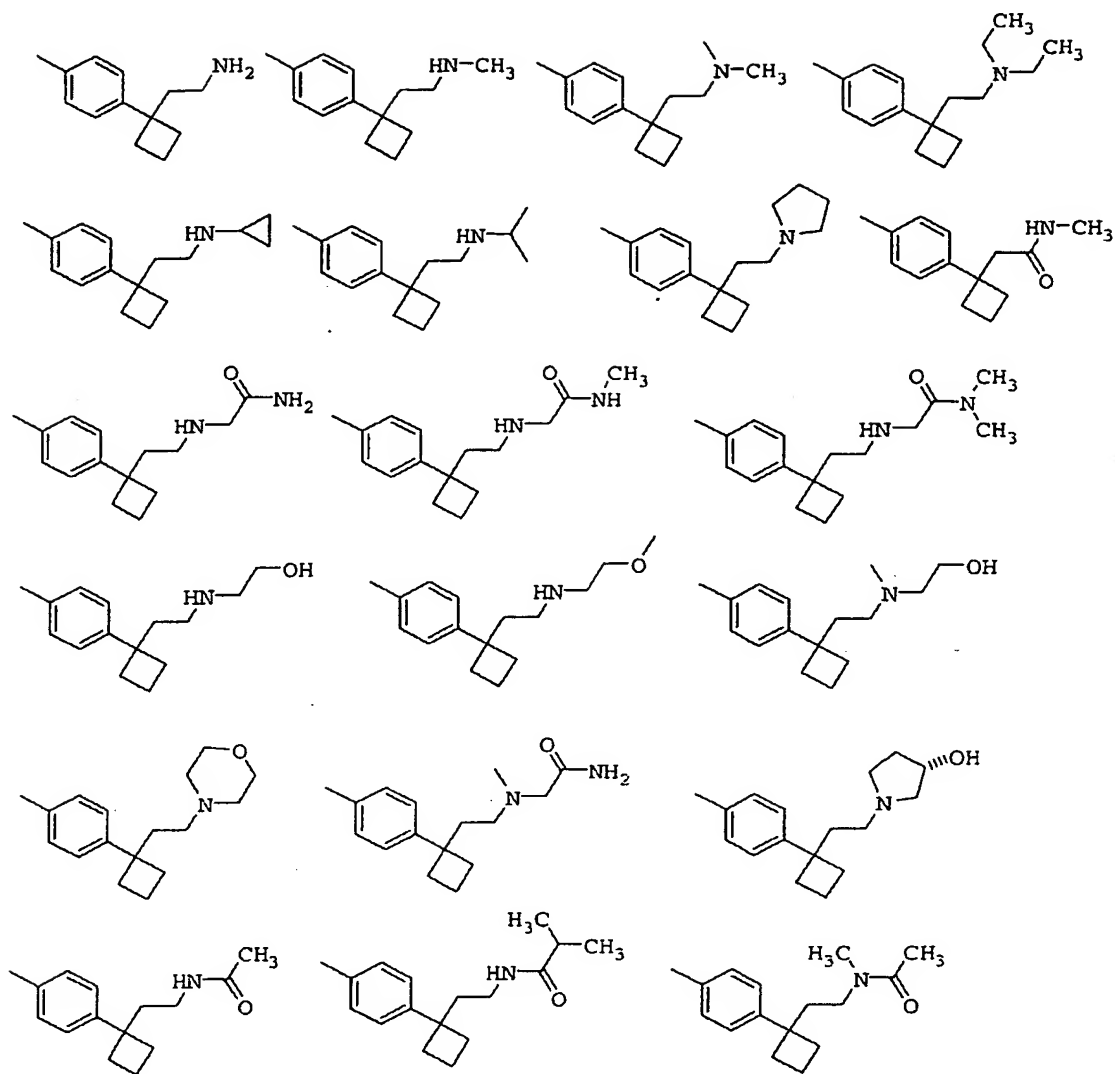


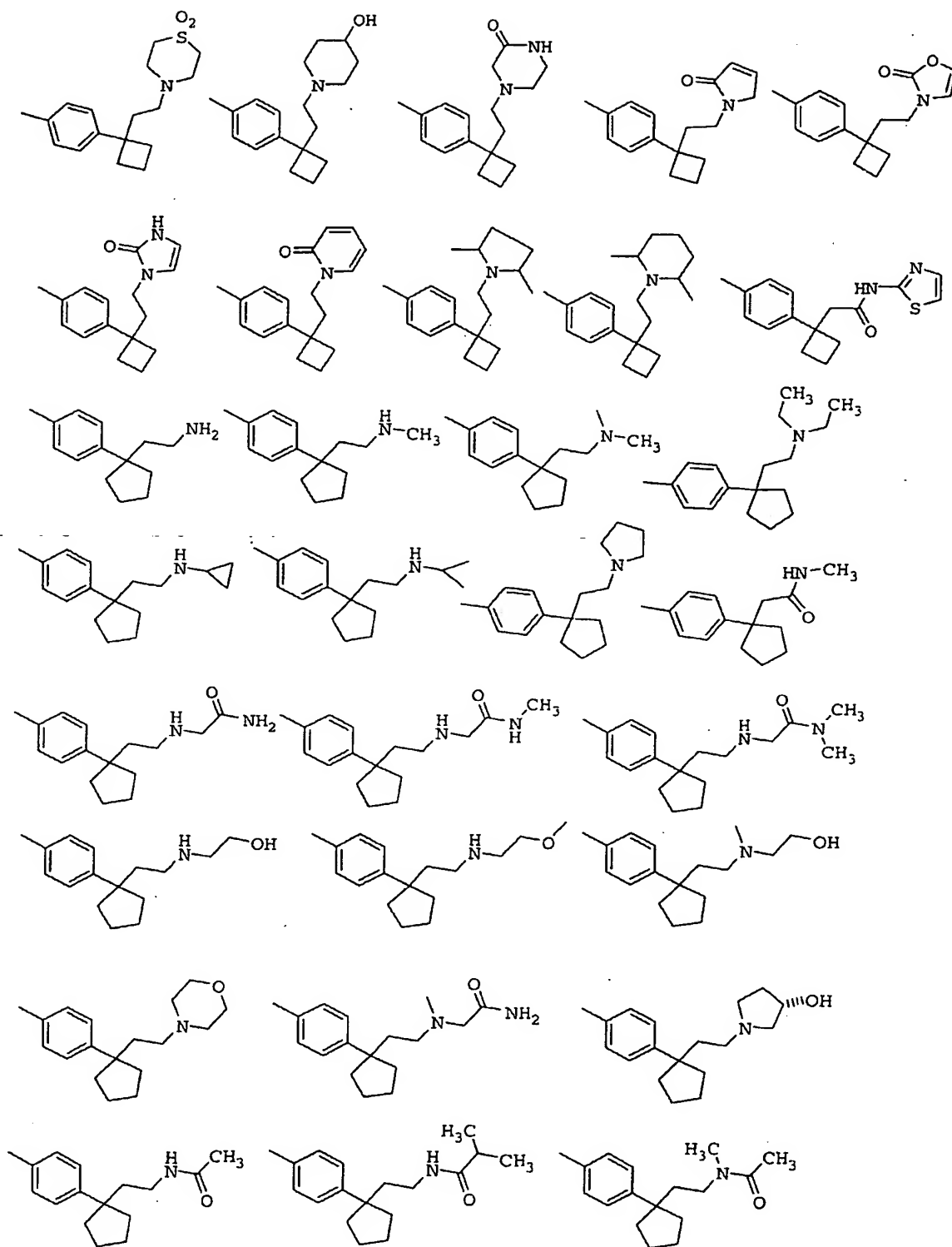


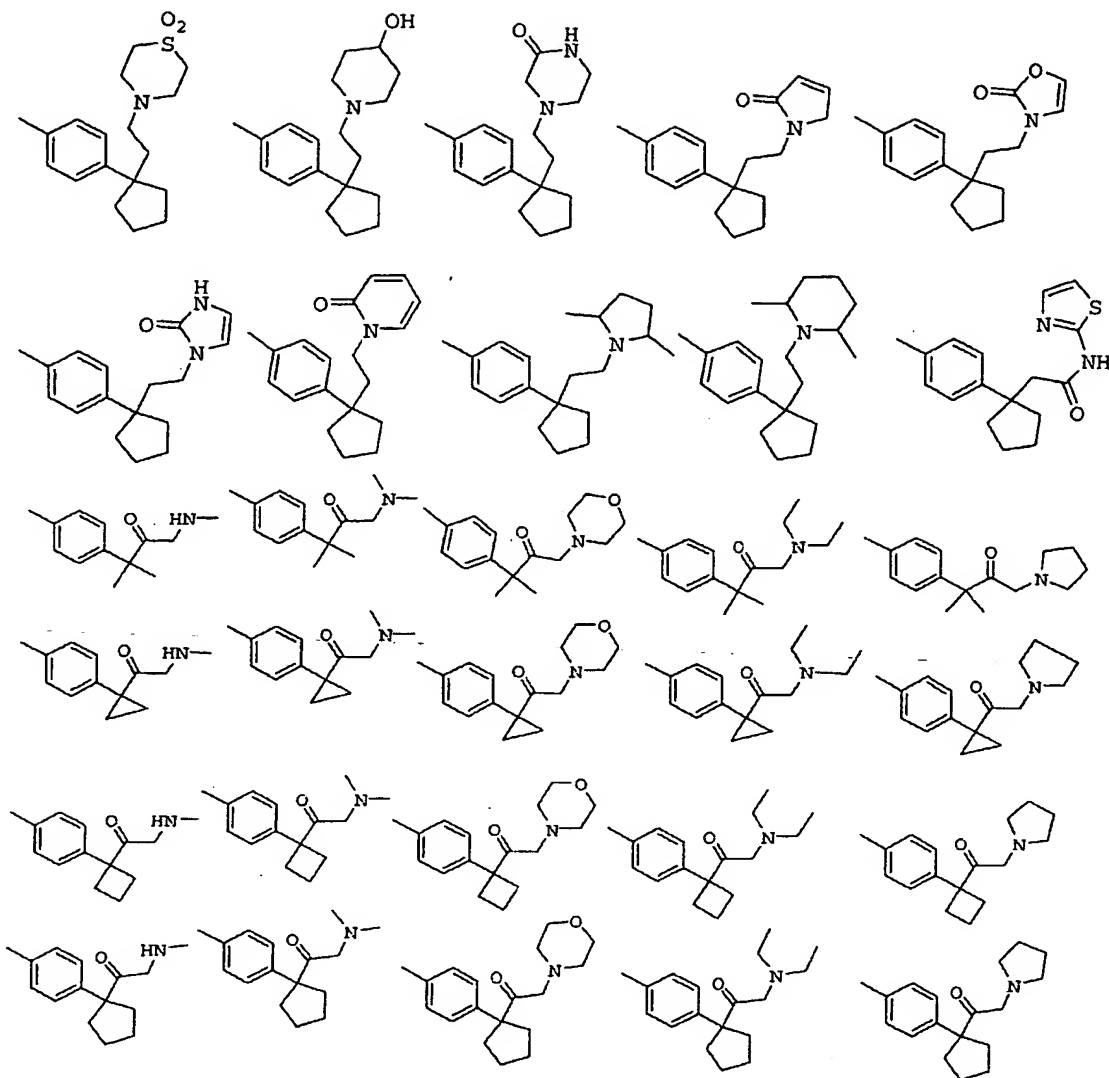












- 5 [15] In another preferred embodiment, the present invention provides a novel compound, wherein the compound is selected from the group:

10 1-[(6-chloro-2-naphthyl)sulfonyl]-4-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl}piperazine;

5-chloro-N-(5-chloro-2-pyridinyl)-2-((4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl)amino)benzamide

;

- N*-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 5 *N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*⁵-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-pyrazole-3,5-dicarboxamide;
- 10 3-cyano-*N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-pyrazole-5-carboxamide;
- N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1*H*-pyrazole-5-carboxamide;
- 15 *N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 20 *N*-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 25 *N*-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 30 1-(2,3-dihydro-1*H*-indol-6-yl)-*N*⁵-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1*H*-pyrazole-3,5-dicarboxamide;
- 35 1-(2,3-dihydro-1*H*-indol-6-yl)-*N*⁵-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1*H*-pyrazole-3,5-dicarboxamide;

5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[2-(dimethylamino)-
1,1-dimethylethyl]benzoyl}amino)benzamide;

5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(methylamino)methyl]cyclopropyl}benzoyl)amino]benzami
de;

10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
(methoxymethyl)cyclopropyl]benzoyl}amino)benzamide;

15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(dimethylamino)methyl]cyclopropyl}benzoyl)amino]benza
mide;

5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-methyl-1-
pyrrolidinyl)methyl]cyclopropyl}benzoyl)amino]benzamid
e;

20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-oxo-1-
pyrrolidinyl)methyl]cyclopropyl}benzoyl)amino]benzamid
e;

25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(isopropylamino)methyl]cyclopropyl}benzoyl)amino]benz
amide;

30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(cyclopropylamino)methyl]cyclopropyl}benzoyl)amino]be
nzamide;

35 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(cyclobutylamino)methyl]cyclopropyl}benzoyl)amino]ben
zamide;

- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((2-hydroxyethyl)amino)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((2-hydroxyethyl)(methyl)amino)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((3-hydroxy-1-pyrrolidinyl)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((4-hydroxy-1-piperidinyl)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((2-oxo-1-piperidinyl)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((2-oxo-1-imidazolidinyl)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-([4-(1-((2-oxo-1-pyrrolidinyl)methyl)cyclopropyl)benzoyl)amino)benzamide;
- 2-([4-(1-([acetyl(methyl)amino)methyl)cyclopropyl)benzoyl)amino)-5-chloro-*N*-(5-chloro-2-pyridinyl)benzamide;

- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
({methyl (methylamino) carbonyl} amino) methyl} cyclopropyl
1] benzyl} amino) benzamide;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-
{ {methyl (methylsulfonyl) amino} methyl} cyclopropyl) benzy
1] amino} benzamide;
- 10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[{methylsulfonyl} amino] cyclopropyl} benzyl) amino] benzam
ide;
- 15 2-((4-[1-(acetylamino) cyclopropyl] benzyl) amino)-5-chloro-*N*-
(5-chloro-2-pyridinyl) benzamide;
- 20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-
hydroxyethyl) amino} methyl} cyclopropyl) benzyl] amino} ben
zamide;
- 25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-
hydroxyethyl) (methyl) amino} methyl} cyclopropyl) benzyl] a
mino} benzamide;
- 30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(1,3-thiazol-2-
ylamino) methyl] cyclopropyl} benzoyl) amino] benzamide;
- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-methyl-1*H*-
imidazol-1-
yl) methyl] cyclopropyl} benzoyl) amino] benzamide;
- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
({{(methylamino) carbonyl} amino) methyl} cyclopropyl] benz
oyl} amino) benzamide;

- methyl [1-(4-{{(4-chloro-2-{{(5-chloro-2-pyridinyl)amino}carbonyl}phenyl)amino}carbonyl}phenyl)cyclopropyl)methylcarbamate;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-
{{(methylsulfonyl)amino}methyl}cyclopropyl)benzoyl}ami
no}benzamide;
- 10 2-{{4-[1-(2-amino-2-oxoethyl)cyclopropyl]benzoyl}amino)-5-
chloro-*N*-(5-chloro-2-pyridinyl)benzamide;
- 15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-[1-[2-
(dimethylamino)-2-
oxoethyl]cyclopropyl}benzyl)amino}benzamide;
- 20 *N*-{4-[1-(2-amino-2-oxoethyl)cyclopropyl]phenyl}-1-(4-
methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*-{4-[1-(aminomethyl)cyclopropyl]phenyl}-1-(4-
methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 25 1-(4-methoxyphenyl)-*N*-(4-[1-
{{(methylamino)methyl}cyclopropyl}phenyl)-1*H*-1,2,3-
triazole-5-carboxamide;
- 30 1-(4-methoxyphenyl)-*N*-{4-[1-(1-
pyrrolidinylmethyl)cyclopropyl]phenyl}-1*H*-1,2,3-
triazole-5-carboxamide;
- 35 1-(4-methoxyphenyl)-*N*⁵-{4-[1-(1-
pyrrolidinylmethyl)cyclopropyl]phenyl}-1*H*-pyrazole-
3,5-dicarboxamide;

- 1- (4-methoxyphenyl) -N⁵- (4- {1- [(2-oxo-1-
pyrrolidinyl)methyl]cyclopropyl}phenyl) -1H-pyrazole-
3,5-dicarboxamide;
- 5 1- (4-methoxyphenyl) -N⁵- (4- {1-
[(methylamino)methyl]cyclopropyl}phenyl) -1H-pyrazole-
3,5-dicarboxamide;
- 10 3-cyano-1- (4-methoxyphenyl) -N- (4- {1-
[(methylamino)methyl]cyclopropyl}phenyl) -1H-pyrazole-
5-carboxamide;
- 15 3-cyano-1- (4-methoxyphenyl) -N- {4- [1- (1-
pyrrolidinylmethyl)cyclopropyl]phenyl} -1H-pyrazole-5-
carboxamide;
- 20 3-cyano-1- (4-methoxyphenyl) -N- (4- {1- [(2-oxo-1-
pyrrolidinyl)methyl]cyclopropyl}phenyl) -1H-pyrazole-5-
carboxamide;
- 1- (4-methoxyphenyl) -3- (methylsulfonyl) -N- (4- {1- [(2-oxo-1-
pyrrolidinyl)methyl]cyclopropyl}phenyl) -1H-pyrazole-5-
carboxamide;
- 25 N- (4- {1- [(3-hydroxy-1-
pyrrolidinyl)methyl]cyclopropyl}phenyl) -1- (4-
methoxyphenyl) -3- (methylsulfonyl) -1H-pyrazole-5-
carboxamide;
- 30 5-chloro-thiophene-2-carboxylic acid {1- [4- (1-pyrrolidin-1-
ylmethyl-cyclopropyl) -benzoyl] -pyrrolidin-3-yl} -amide
;
- 35 5-chloro-thiophene-2-carboxylic acid {1- [4- (1-
dimethylaminomethyl-cyclopropyl) -benzoyl] -pyrrolidin-
3-yl} -amide;

- 3-chloro-1H-indole-6-carboxylic acid {1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid {1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 15 2{4-[4-chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;
- 20 2{4-[4-chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propyl alcohol;
- 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-[2-(ethylamino)-1,1-dimethylethyl]benzoyl}amino)benzamide;
- 25 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl}amino)benzamide;
- 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl}amino)benzamide;
- 30 2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;
- 35 2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-4-methoxy-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;

N-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)benzamide;

5 *N*-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)-5-methoxybenzamide;

N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)benzamide;

10

N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl]amino)benzamide;

15 *N*-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)-5-methoxybenzamide;

2-[4-(2-[acetyl(methyl)amino]-1,1-dimethylethyl)benzoyl]amino]-*N*-(5-chloropyridin-2-yl)benzamide;

20

2-(4-([2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylamino]methyl)-phenyl)-2-methyl-propionic acid methyl ester;

25 5-chloro-*N*-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)benzamide;

5-chloro-*N*-(5-chloro-pyridin-2-yl)-2-[4-(2-dimethylamino-1,1-dimethyl-ethyl)-benzylamino]-benzamide;

30

N-(5-chloropyridin-2-yl)-2-([4-[1-(hydroxymethyl)cyclopropyl]benzoyl]amino)-5-methoxybenzamide;

35 *N*-(5-chloropyridin-2-yl)-5-methoxy-2-([4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl]amino)benzamide;

- N*-(5-chloropyridin-2-yl)-2-((4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl)amino)benzamide;
- 5 1-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-cyclopropanecarboxylic acid methyl ester;
- N*-(5-chloropyridin-2-yl)-2-((4-[1-(hydroxymethyl)cyclopropyl]benzoyl)amino)benzamide;
- 10 6-chloro-3-(5-chloropyridin-2-yl)-2-[4-(1,1-dimethyl-2-morpholin-4-ylethyl)phenyl]quinazolin-4(3*H*)-one;
- 15 3-(5-chloropyridin-2-yl)-2-{4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]phenyl}quinazolin-4(3*H*)-one;
- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 20 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 25 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-amide;
- 30 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid [4-(1-{2-[(2-hydroxy-ethyl)-methylamino]-ethyl}-cyclopropyl)-phenyl]-amide;
- 35 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid (4-{1-[2-(carbamoylmethyl-methylamino)-ethyl]-cyclopropyl}-phenyl)-amide;

- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-amide;
- 5 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 10 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 15 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclobutyl)-phenyl]-amide;
- 20 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-carbamoylmethyl-cyclobutyl)-phenyl]-amide;
- 25 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-methylamino-ethyl)-cyclobutyl]-phenyl}-amide;
- 30 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-amide;
- 35 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclobutyl]-phenyl}-amide;
- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-morpholin-4-yl-ethyl)-cyclobutyl]-phenyl}-amide;

- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-amide;
- 5 5-cyano-2-(4-methoxy-phenyl)-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 10 2-(4-methoxy-phenyl)-5-methyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 15 1-(4-methoxy-phenyl)-1H-pyrazole-3,5-dicarboxylic acid 3-amide 5-({4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide);
- 20 5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 25 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid [4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 30 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 2-[1-(4-{2-[3-(4-methoxy-phenyl)-3H-[1,2,3]triazol-4-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-N-methyl-acetamide;
- 35 2-[1-(4-{2-[3-(4-methoxy-phenyl)-3H-[1,2,3]triazol-4-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-acetamide;

- 2-[1-(4-(2-[2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazol-3-yl]-2-oxo-ethyl)-phenyl)-cyclopropyl]-acetamide;
- 5 2-[1-(4-(2-[5-cyano-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl)-phenyl)-cyclopropyl]-acetamide;
- 10 2-[1-(4-(2-[5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl)-phenyl)-cyclopropyl]-acetamide;
- 15 2-[1-(4-(2-[5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl)-phenyl)-cyclopropyl]-N-methyl-acetamide;
- 20 5-chloro-N-(5-chloro-2-pyridinyl)-2-((4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl)amino)benzamide;
- N-(5-chloro-2-pyridinyl)-5-methoxy-2-((4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl)amino)benzamide;
- 25 N-(5-chloro-2-pyridinyl)-5-fluoro-2-((4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl)amino)benzamide;
- 30 N-(5-chloro-2-pyridinyl)-5-methyl-2-((4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl)amino)benzamide;
- 35 N-(5-chloro-2-pyridinyl)-5-methylsulfonyl-2-((4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl)amino)benzamide;

N-(5-chloro-2-pyridinyl)-5-cyano-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

5 N-(5-chloro-2-pyridinyl)-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-pyridine-2-carboxylic acid (5-chloro-pyridin-2-yl)-amide;

10 N-(5-chloro-pyridin-2-yl)-4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-nicotinamide;

15 N-(5-chloro-pyridin-2-yl)-3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-isonicotinamide;

N-(5-chloro-pyridin-2-yl)-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-nicotinamide;

20 5-chloro-N-(5-chloro-2-pyridinyl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

25 N-(5-chloro-2-pyridinyl)-5-methoxy-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

30 N-(5-chloro-2-pyridinyl)-5-fluoro-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

35 N-(5-chloro-2-pyridinyl)-5-methyl-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

- N-(5-chloro-2-pyridinyl)-5-methylsulfonyl-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}benzoylamino)benzamide;
- 5 N-(5-chloro-2-pyridinyl)-5-cyano-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;
- 10 N-(5-chloro-2-pyridinyl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;
- 3-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-pyridine-2-carboxylic acid (5-chloro-pyridin-2-yl)-amide;
- 15 N-(5-chloro-pyridin-2-yl)-4-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-nicotinamide;
- N-(5-chloro-pyridin-2-yl)-3-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-isonicotinamide;
- 20 N-(5-chloro-pyridin-2-yl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-nicotinamide;
- 25 3-chloro-1H-indole-6-carboxylic acid {4-dimethylcarbamoyl-2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid {5-dimethylcarbamoyl-2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 35 3-chloro-1H-indole-6-carboxylic acid {4-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-tetrahydro-pyran-3-yl}-amide;

- 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-tetrahydro-pyran-4-yl}-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid (1,1-dioxo-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-hexahydro-1 λ ⁶-thiopyran-4-yl)-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid (1,1-dioxo-4-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-hexahydro-1 λ ⁶-thiopyran-3-yl)-amide;
- 15 3-chloro-1H-indole-6-carboxylic acid (1-acetyl-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl)-amide;
- 20 3-chloro-1H-indole-6-carboxylic acid (1-acetyl-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl)-amide;
- 4-[(3-chloro-1H-indole-6-carbonyl)-amino]-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidine-1-carboxylic acid methyl ester;
- 25 3-chloro-1H-indole-6-carboxylic acid (1-(2-methoxy-acetyl)-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid (2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-cyclopentyl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid (4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-tetrahydro-furan-3-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid {1-acetyl-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidin-3-yl}-amide;
- 5 5-chloro-thiophene-2-carboxylic acid {1-cyclopropanecarbonyl-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidin-3-yl}-amide;
- 10 3-[(5-chloro-thiophene-2-carbonyl)-amino]-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidine-1-carboxylic acid methyl ester;
- 15 5-chloro-thiophene-2-carboxylic acid [4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1-(2-methoxy-acetyl)-pyrrolidin-3-yl]-amide;
- 20 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-4-dimethylcarbamoyl-cyclopentyl}-amide;
- 25 5-chloro-thiophene-2-carboxylic acid {1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-indan-2-yl}-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1,2,3,4-tetrahydro-naphthalen-2-yl}-amide;
- 35 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-7-oxa-bicyclo[2.2.1]hept-2-yl}-amide;
- 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-4-dimethylcarbamoyl-cyclopentyl}-amide;

- 5-chloro-thiophene-2-carboxylic acid (8-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1-oxa-spiro[4.4]non-7-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (8-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1-oxa-spiro[4.4]non-7-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-cyclopentyl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-4-dimethylcarbamoyl-cyclopentyl)-amide;
- 20 3-[(5-chloro-thiophene-2-carbonyl)-amino]-4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-pyrrolidine-1-carboxylic acid methyl ester;
- 25 5-chloro-thiophene-2-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-furan-3-yl)-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-cyclohexyl)-amide;
- 35 3-chloro-1H-indole-6-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-4-dimethylcarbamoyl-cyclohexyl)-amide;
- 4-[(3-Chloro-1H-indole-6-carbonyl)-amino]-3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-piperidine-1-carboxylic acid methyl ester;

- 3-chloro-1H-indole-6-carboxylic acid (3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1,1-dioxo-hexahydro-1 λ ⁶-thiopyran-4-yl)-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1,1-dioxo-hexahydro-1 λ ⁶-thiopyran-3-yl)-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-pyran-3-yl)-amide;
- 15 3-chloro-1H-indole-6-carboxylic acid (3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-pyran-4-yl)-amide;
- (1R, 2S)-5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide;
- 20 (1R, 2S)-3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide;
- 25 (1R, 2S)-5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide; and,
- 30 Cis-3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-phenylcarbamoyl]-cyclohexyl}-amide;

or a pharmaceutically acceptable salt form thereof.

35

In another embodiment, the present invention provides novel pharmaceutical compositions, comprising: a

pharmaceutically acceptable carrier and a therapeutically effective amount of a compound of the present invention or a pharmaceutically acceptable salt form thereof.

5

In another embodiment, the present invention provides a novel method for treating a thromboembolic disorder, comprising: administering to a patient in need thereof a therapeutically effective amount of a compound of the present invention or a pharmaceutically acceptable salt form thereof.

In another preferred embodiment, the present invention provides a novel method, wherein the thromboembolic disorder is selected from the group consisting of arterial cardiovascular thromboembolic disorders, venous cardiovascular thromboembolic disorders, and thromboembolic disorders in the chambers of the heart.

20

In another preferred embodiment, the present invention provides a novel method, wherein the thromboembolic disorder is selected from unstable angina, an acute coronary syndrome, first myocardial infarction, recurrent myocardial infarction, ischemic sudden death, transient ischemic attack, stroke, atherosclerosis, peripheral occlusive arterial disease, venous thrombosis, deep vein thrombosis, thrombophlebitis, arterial embolism, coronary arterial thrombosis, cerebral arterial thrombosis, cerebral embolism, kidney embolism, pulmonary embolism, and thrombosis resulting from (a) prosthetic valves or other implants, (b) indwelling catheters, (c) stents, (d) cardiopulmonary bypass, (e) hemodialysis, or (f) other procedures in which blood is exposed to an artificial surface that promotes thrombosis.

35

In another embodiment, the present invention provides a novel method of treating a patient in need of
5 thromboembolic disorder treatment, comprising:
administering a compound of the present invention or a
pharmaceutically acceptable salt form thereof in an amount
effective to treat a thromboembolic disorder

10

In another embodiment, the present invention provides a novel method, comprising: administering a compound of the present invention or a pharmaceutically acceptable salt form thereof in an amount effective to treat a
15 thromboembolic disorder.

In another embodiment, the present invention provides a novel method for treating a thromboembolic disorder,
20 comprising: administering to a patient in need thereof a therapeutically effective amount of a first and second therapeutic agent, wherein the first therapeutic agent is compound of the present invention or a pharmaceutically acceptable salt thereof and the second therapeutic agent is
25 at least one agent selected from a second factor Xa inhibitor, an anti-coagulant agent, an anti-platelet agent, a thrombin inhibiting agent, a thrombolytic agent, and a fibrinolytic agent.

30

In another preferred embodiment, the present invention provides a novel method, wherein the second therapeutic agent is at least one agent selected from warfarin, unfractionated heparin, low molecular weight heparin,
35 synthetic pentasaccharide, hirudin, argatrobanas, aspirin, ibuprofen, naproxen, sulindac, indomethacin, mefenamate,

droxicam, diclofenac, sulfinpyrazone, piroxicam,
ticlopidine, clopidogrel, tirofiban, eptifibatide,
abciximab, melagatran, disulfatohirudin, tissue plasminogen
activator, modified tissue plasminogen activator,
5 anistreplase, urokinase, and streptokinase.

In another preferred embodiment, the present invention
provides a novel method, wherein the second therapeutic
10 agent is at least one anti-platelet agent.

In another preferred embodiment, the present invention
provides a novel method, wherein the anti-platelet agent is
15 aspirin and clopidogrel.

In another preferred embodiment, the present invention
provides a novel method, wherein the anti-platelet agent is
20 clopidogrel.

In another embodiment, the present invention provides
a novel article of manufacture, comprising:

- 25 (a) a first container;
(b) a pharmaceutical composition located within the
first container, wherein the composition, comprises: a
first therapeutic agent, comprising: a compound of the
present invention or a pharmaceutically acceptable salt
30 form thereof; and,
(c) a package insert stating that the pharmaceutical
composition can be used for the treatment of a
thromboembolic disorder.

35

In another preferred embodiment, the present invention provides a novel article of manufacture, further comprising:

- (d) a second container;
- 5 wherein components (a) and (b) are located within the second container and component (c) is located within or outside of the second container.

10 In another embodiment, the present invention provides a novel article of manufacture, comprising:

- (a) a first container;
- (b) a pharmaceutical composition located within the first container, wherein the composition, comprises: a
15 first therapeutic agent, comprising: a compound of the present invention or a pharmaceutically acceptable salt form thereof; and,
- (c) a package insert stating that the pharmaceutical composition can be used in combination with a second
20 therapeutic agent to treat a thromboembolic disorder.

In another preferred embodiment, the present invention provides a novel article of manufacture, further
25 comprising:

- (d) a second container;
- wherein components (a) and (b) are located within the second container and component (c) is located within or outside of the second container.

30

In another embodiment, the present invention provides a compound of the present invention for use in therapy.

35

In another embodiment, the present invention provides the use of a compound of the present invention as described above for the manufacture of a medicament for the treatment of a thromboembolic disorder.

5

The present invention may be embodied in other specific forms without departing from the spirit or essential attributes thereof. This invention encompasses all combinations of preferred aspects of the invention
10 noted herein. It is understood that any and all embodiments of the present invention may be taken in conjunction with any other embodiment or embodiments to describe additional more preferred embodiments. It is also to be understood that each individual element of the
15 preferred embodiments is its own independent preferred embodiment. Furthermore, any element of an embodiment is meant to be combined with any and all other elements from any embodiment to describe an additional embodiment.

20

DEFINITIONS

The compounds herein described may have asymmetric centers. Compounds of the present invention containing an asymmetrically substituted atom may be isolated in optically active or racemic forms. It is well known in the
25 art how to prepare optically active forms, such as by resolution of racemic forms or by synthesis from optically active starting materials. Many geometric isomers of olefins, C=N double bonds, and the like can also be present in the compounds described herein, and all such stable
30 isomers are contemplated in the present invention. Cis and trans geometric isomers of the compounds of the present invention are described and may be isolated as a mixture of isomers or as separated isomeric forms. All chiral, diastereomeric, racemic forms and all geometric isomeric
35 forms of a structure are intended, unless the specific stereochemistry or isomeric form is specifically indicated.

All processes used to prepare compounds of the present invention and intermediates made therein are considered to be part of the present invention. All tautomers of shown or described compounds are also considered to be part of the present invention.

Preferably, the molecular weight of compounds of the present invention is less than about 500, 550, 600, 650, 700, 750, or 800 grams per mole. Preferably, the molecular weight is less than about 800 grams per mole. More preferably, the molecular weight is less than about 750 grams per mole. Even more preferably, the molecular weight is less than about 700 grams per mole.

The term "substituted," as used herein, means that any one or more hydrogens on the designated atom is replaced with a selection from the indicated group, provided that the designated atom's normal valency is not exceeded, and that the substitution results in a stable compound. When a substituent is keto (i.e., =O), then 2 hydrogens on the atom are replaced. Keto substituents are not present on aromatic moieties. Ring double bonds, as used herein, are double bonds that are formed between two adjacent ring atoms (e.g., C=C, C=N, or N=N). The present invention, in general, does not cover groups such as N-halo, S(O)H, and SO₂H.

The present invention is intended to include all isotopes of atoms occurring in the present compounds. Isotopes include those atoms having the same atomic number but different mass numbers. By way of general example and without limitation, isotopes of hydrogen include tritium and deuterium. Isotopes of carbon include C-13 and C-14.

When any variable (e.g., R⁶) occurs more than one time in any constituent or formula for a compound, its definition at each occurrence is independent of its definition at every other occurrence. Thus, for example, if a group is shown to be substituted with 0-2 R⁶, then said group may optionally be substituted with up to two R⁶

groups and R⁶ at each occurrence is selected independently from the definition of R⁶. Also, combinations of substituents and/or variables are permissible only if such combinations result in stable compounds.

5 When a bond to a substituent is shown to cross a bond connecting two atoms in a ring, then such substituent may be bonded to any atom on the ring. When a substituent is listed without indicating the atom via which such substituent is bonded to the rest of the compound of a
10 given formula, then such substituent may be bonded via any atom in such substituent. Combinations of substituents and/or variables are permissible only if such combinations result in stable compounds.

 In cases wherein there are nitrogen atoms (e.g.,
15 amines) on compounds of the present invention, these can be converted to N-oxides by treatment with an oxidizing agent (e.g., MCPBA and/or hydrogen peroxides) to afford other compounds of this invention. Thus, all shown and claimed nitrogen atoms are considered to cover both the shown
20 nitrogen and its N-oxide (N→O) derivative.

 As used herein, "alkyl" is intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms. C₁₋₆ alkyl, is intended to include C₁, C₂, C₃, C₄, C₅, and C₆
25 alkyl groups. Examples of alkyl include, but are not limited to, methyl, ethyl, n-propyl, i-propyl, n-butyl, s-butyl, t-butyl, n-pentyl, and s-pentyl. "Haloalkyl" is intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified
30 number of carbon atoms, substituted with 1 or more halogen (for example -C_vF_w where v = 1 to 3 and w = 1 to (2v+1)). Examples of haloalkyl include, but are not limited to, trifluoromethyl, trichloromethyl, pentafluoroethyl, and pentachloroethyl. "Alkoxy" represents an alkyl group as
35 defined above with the indicated number of carbon atoms

attached through an oxygen bridge. C₁₋₆ alkoxy, is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ alkoxy groups. Examples of alkoxy include, but are not limited to, methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, s-butoxy, t-butoxy, n-pentoxo, and s-pentoxo. "Cycloalkyl" is intended to include saturated ring groups, such as cyclopropyl, cyclobutyl, or cyclopentyl. C₃₋₇ cycloalkyl is intended to include C₃, C₄, C₅, C₆, and C₇ cycloalkyl groups. Alkenyl" is intended to include hydrocarbon chains of either straight or branched configuration and one or more unsaturated carbon-carbon bonds that may occur in any stable point along the chain, such as ethenyl and propenyl. C₂₋₆ alkenyl is intended to include C₂, C₃, C₄, C₅, and C₆ alkenyl groups. "Alkynyl" is intended to include hydrocarbon chains of either straight or branched configuration and one or more triple carbon-carbon bonds that may occur in any stable point along the chain, such as ethynyl and propynyl. C₂₋₆ Alkynyl is intended to include C₂, C₃, C₄, C₅, and C₆ alkynyl groups.

"Halo" or "halogen" as used herein refers to fluoro, chloro, bromo, and iodo; and "counterion" is used to represent a small, negatively charged species such as chloride, bromide, hydroxide, acetate, and sulfate.

As used herein, "carbocycle" or "carbocyclic residue" is intended to mean any stable 3, 4, 5, 6, or 7-membered monocyclic or bicyclic or 7, 8, 9, 10, 11, 12, or 13-membered bicyclic or tricyclic ring, any of which may be saturated, partially unsaturated, or unsaturated (aromatic). Examples of such carbocycles include, but are not limited to, cyclopropyl, cyclobutyl, cyclobutenyl, cyclopentyl, cyclopentenyl, cyclohexyl, cycloheptenyl, cycloheptyl, cycloheptenyl, adamantyl, cyclooctyl, cyclooctenyl, cyclooctadienyl, [3.3.0]bicyclooctane, [4.3.0]bicyclononane, [4.4.0]bicyclodecane, [2.2.2]bicyclooctane, fluorenyl, phenyl, naphthyl, indanyl, adamantyl, and tetrahydronaphthyl. As shown above, bridged

rings are also included in the definition of carbocycle (e.g., [2.2.2]bicyclooctane). A bridged ring occurs when one or more carbon atoms link two non-adjacent carbon atoms. Preferred bridges are one or two carbon atoms. It is noted that a bridge always converts a monocyclic ring into a tricyclic ring. When a ring is bridged, the substituents recited for the ring may also be present on the bridge.

As used herein, the term "heterocycle" or "heterocyclic group" is intended to mean a stable 5, 6, or 7-membered monocyclic or bicyclic or 7, 8, 9, or 10-membered bicyclic heterocyclic ring which is saturated, partially unsaturated or unsaturated (aromatic), and which consists of carbon atoms and 1, 2, 3, or 4 ring heteroatoms independently selected from the group consisting of N, O and S and including any bicyclic group in which any of the above-defined heterocyclic rings is fused to a benzene ring. The nitrogen and sulfur heteroatoms may optionally be oxidized (i.e., N=O and S(O)_p). The nitrogen atom may be substituted or unsubstituted (i.e., N or NR wherein R is H or another substituent, if defined). The heterocyclic ring may be attached to its pendant group at any heteroatom or carbon atom that results in a stable structure. The heterocyclic rings described herein may be substituted on carbon or on a nitrogen atom if the resulting compound is stable. A nitrogen in the heterocycle may optionally be quaternized. It is preferred that when the total number of S and O atoms in the heterocycle exceeds 1, then these heteroatoms are not adjacent to one another. It is preferred that the total number of S and O atoms in the heterocycle is not more than 1. As used herein, the term "aromatic heterocyclic group" or "heteroaryl" is intended to mean a stable 5, 6, or 7-membered monocyclic or bicyclic or 7, 8, 9, or 10-membered bicyclic heterocyclic aromatic ring which consists of carbon atoms and 1, 2, 3, or 4 heterotams independently selected from the group consisting

of N, O and S. The nitrogen atom may be substituted or unsubstituted (i.e., N or NR wherein R is H or another substituent, if defined). The nitrogen and sulfur heteroatoms may optionally be oxidized (i.e., N=O and S(O)_p). It is to be noted that total number of S and O atoms in the aromatic heterocycle is not more than 1. Bridged rings are also included in the definition of heterocycle. A bridged ring occurs when one or more atoms (i.e., C, O, N, or S) link two non-adjacent carbon or nitrogen atoms. Preferred bridges include, but are not limited to, one carbon atom, two carbon atoms, one nitrogen atom, two nitrogen atoms, and a carbon-nitrogen group. It is noted that a bridge always converts a monocyclic ring into a tricyclic ring. When a ring is bridged, the substituents recited for the ring may also be present on the bridge.

Examples of heterocycles include, but are not limited to, acridinyl, azocinyl, benzimidazolyl, benzofuranyl, benzothiofuranyl, benzothiophenyl, benzoxazolyl, benzoxazolinyll, benzthiazolyl, benztriazolyl, benztetrazolyl, benzisoxazolyl, benzisothiazolyl, benzimidazolinyll, carbazolyl, 4aH-carbazolyl, carbolinyl, chromanyl, chromenyl, cinnolinyl, decahydroquinolinyl, 2H,6H-1,5,2-dithiazinyl, dihydrofuro[2,3-b]tetrahydrofuran, furanyl, furazanyl, imidazolidinyl, imidazolinyll, imidazolyl, 1H-indazolyl, indolenyl, indolinyl, indolizinyll, indolyl, 3H-indolyl, isatinoyl, isobenzofuranyl, isochromanyl, isoindazolyl, isoindolinyl, isoindolyl, isoquinolinyl, isothiazolyl, isoxazolyl, methylenedioxyphenyl, morpholinyl, naphthyridinyl, octahydroisoquinolinyl, oxadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolidinyl, oxazolyl, oxindolyl, pyrimidinyl, phenanthridinyl, phenanthrolinyl, phenazinyl, phenothiazinyl, phenoxathinyl, phenoxazinyl, phthalazinyl, piperazinyl, piperidinyl, piperidonyl, 4-piperidonyl,

piperonyl, pteridinyl, purinyl, pyranyl, pyrazinyl, pyrazolidinyl, pyrazolinyl, pyrazolyl, pyridazinyl, pyridooxazole, pyridoimidazole, pyridothiazole, pyridinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolinyl, 5 2H-pyrrolyl, pyrrolyl, quinazolinyl, quinolinyl, 4H-quinoliziny, quinoxaliny, quinuclidiny, tetrahydrofurany, tetrahydroisoquinolinyl, tetrahydroquinolinyl, tetrazolyl, 6H-1,2,5-thiadiazinyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 10 1,3,4-thiadiazolyl, thianthrenyl, thiazolyl, thienyl, thienothiazolyl, thienooxazolyl, thienoimidazolyl, thiophenyl, triazinyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl, 1,3,4-triazolyl, and xanthenyl. Also included are fused ring and spiro compounds containing, for 15 example, the above heterocycles.

The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in 20 contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

As used herein, "pharmaceutically acceptable salts" 25 refer to derivatives of the disclosed compounds wherein the parent compound is modified by making acid or base salts thereof. Examples of pharmaceutically acceptable salts include, but are not limited to, mineral or organic acid salts of basic residues such as amines; alkali or organic 30 salts of acidic residues such as carboxylic acids; and the like. The pharmaceutically acceptable salts include the conventional non-toxic salts or the quaternary ammonium salts of the parent compound formed, for example, from non-toxic inorganic or organic acids. For example, such 35 conventional non-toxic salts include, but are not limited to, those derived from inorganic and organic acids selected

from 2-acetoxybenzoic, 2-hydroxyethane sulfonic, acetic, ascorbic, benzene sulfonic, benzoic, bicarbonic, carbonic, citric, edetic, ethane disulfonic, ethane sulfonic, fumaric, glucoheptonic, gluconic, glutamic, glycolic, glycollyarsanilic, hexylresorcinic, hydrabamic, hydrobromic, hydrochloric, hydroiodide, hydroxymaleic, hydroxynaphthoic, isethionic, lactic, lactobionic, lauryl sulfonic, maleic, malic, mandelic, methane sulfonic, napsylic, nitric, oxalic, pamoic, pantothenic, phenylacetic, phosphoric, polygalacturonic, propionic, salicyclic, stearic, subacetic, succinic, sulfamic, sulfanilic, sulfuric, tannic, tartaric, and toluene sulfonic.

The pharmaceutically acceptable salts of the present invention can be synthesized from the parent compound that contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in *Remington's Pharmaceutical Sciences*, 18th ed., Mack Publishing Company, Easton, PA, 1990, p 1445, the disclosure of which is hereby incorporated by reference.

Since prodrugs are known to enhance numerous desirable qualities of pharmaceuticals (e.g., solubility, bioavailability, manufacturing, etc.) the compounds of the present invention may be delivered in prodrug form. Thus, the present invention is intended to cover prodrugs of the presently claimed compounds, methods of delivering the same and compositions containing the same. "Prodrugs" are intended to include any covalently bonded carriers that release an active parent drug of the present invention in vivo when such prodrug is administered to a mammalian

subject. Prodrugs the present invention are prepared by modifying functional groups present in the compound in such a way that the modifications are cleaved, either in routine manipulation or *in vivo*, to the parent compound. Prodrugs
5 include compounds of the present invention wherein a hydroxy, amino, or sulfhydryl group is bonded to any group that, when the prodrug of the present invention is administered to a mammalian subject, it cleaves to form a free hydroxyl, free amino, or free sulfhydryl group,
10 respectively. Examples of prodrugs include, but are not limited to, acetate, formate, and benzoate derivatives of alcohol and amine functional groups in the compounds of the present invention.

"Stable compound" and "stable structure" are meant to
15 indicate a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into an efficacious therapeutic agent. It is preferred that there presently recited compounds do not contain a N-halo, $S(O)_2H$, or $S(O)H$ group.

20 "Substituted" is intended to indicate that one or more hydrogens on the atom indicated in the expression using "substituted" is replaced with a selection from the indicated group(s), provided that the indicated atom's normal valency is not exceeded, and that the substitution
25 results in a stable compound. When a substituent is keto (i.e., $=O$) group, then 2 hydrogens on the atom are replaced.

As used herein, "treating" or "treatment" cover the treatment of a disease-state in a mammal, particularly in a
30 human, and include: (a) preventing the disease-state from occurring in a mammal, in particular, when such mammal is predisposed to the disease-state but has not yet been diagnosed as having it; (b) inhibiting the disease-state, i.e., arresting its development; and/or (c) relieving the
35 disease-state, i.e., causing regression of the disease state.

"Therapeutically effective amount" is intended to include an amount of a compound of the present invention that is effective when administered alone or in combination to inhibit factor Xa. "Therapeutically effective amount" is also intended to include an amount of the combination of compounds claimed that is effective to inhibit factor Xa. The combination of compounds is preferably a synergistic combination. Synergy, as described, for example, by Chou and Talalay, *Adv. Enzyme Regul.* **1984**, 22:27-55, occurs when the effect (in this case, inhibition of factor Xa) of the compounds when administered in combination is greater than the additive effect of the compounds when administered alone as a single agent. In general, a synergistic effect is most clearly demonstrated at sub-optimal concentrations of the compounds. Synergy can be in terms of lower cytotoxicity, increased antithrombotic effect, or some other beneficial effect of the combination compared with the individual components.

20 SYNTHESIS

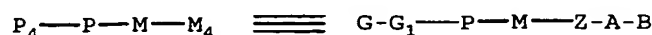
The compounds of the present invention can be prepared in a number of ways known to one skilled in the art of organic synthesis. The compounds of the present invention can be synthesized using the methods described below, together with synthetic methods known in the art of synthetic organic chemistry, or by variations thereon as appreciated by those skilled in the art. Preferred methods include, but are not limited to, those described below. The reactions are performed in a solvent appropriate to the reagents and materials employed and suitable for the transformations being effected. It will be understood by those skilled in the art of organic synthesis that the functionality present on the molecule should be consistent with the transformations proposed. This will sometimes require a judgment to modify the order of the synthetic steps or to select one particular process scheme over

another in order to obtain a desired compound of the invention.

It will also be recognized that another major consideration in the planning of any synthetic route in this field is the judicious choice of the protecting group used for protection of the reactive functional groups present in the compounds described in this invention. An authoritative account describing the many alternatives to the trained practitioner is Greene and Wuts (*Protective Groups In Organic Synthesis*, Wiley and Sons, 1991). All references cited herein are hereby incorporated in their entirety herein by reference.

The compounds of the present invention of formula I (Scheme 1) where P is not fused onto ring M can be prepared as outlined in Scheme 2 to Scheme 10 and via standard methods known to those skilled in the art.

Scheme 1



Formula I

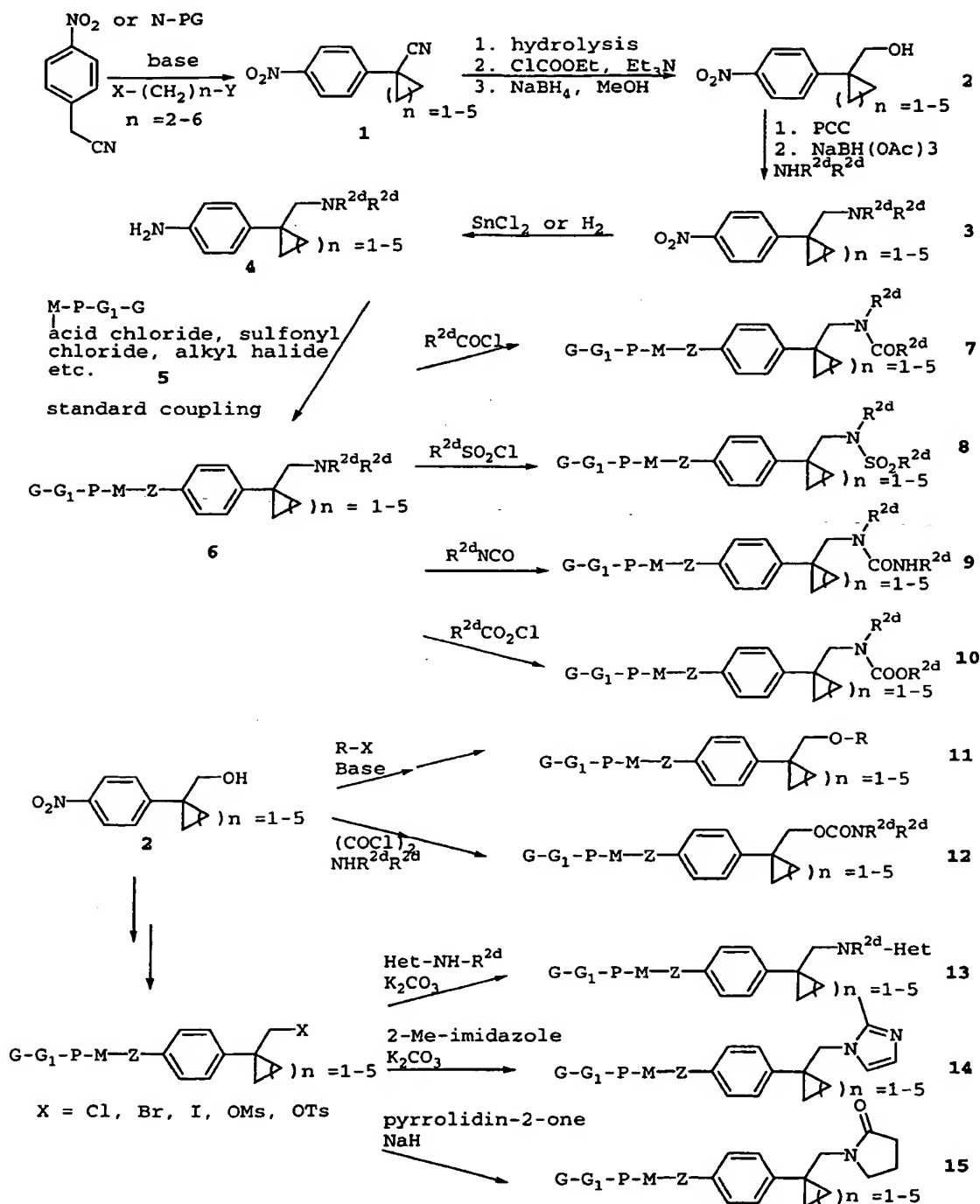
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The compounds of the present invention of formula I where Y is C₃-C₇ cycloalkyl can be prepared as shown in Scheme 2. Commercially available 4-nitrophenylacetonitrile (or properly protected 4-aminophenylacetonitrile) can be used as the starting material. Alkylation with NaH, KtOBu, NaNH₂, n-BuLi, s-BuLi, NaOEt, or aq NaOH, etc. as the base, and X-(CH₂)_n-Y (X, Y can be Cl, Br, I, OMs or OTs, ⁺S(CH₃)₂, n = 2-6) as the alkylating reagent can afford the cycloalkyl intermediate 1. Hydrolysis of the nitrile group, followed by reduction of the ester group can provide the alcohol 2. Oxidation of 2, then reductive amination with NHR^{2d}R^{2d} will provide 3. Reduction of the nitro group or deprotection of the amino group can produce the A-B precursor 4, which can be coupled with 5 using standard

coupling conditions to provide **6**. When one of the R^{2d} groups is H, **6** can react with acid chlorides, carbamoyl chlorides, sulfonyl chlorides, and isocyanates to provide compounds of the invention with structures **7**, **8**, **9**, and **10**.

- 5 Alternatively, alcohol **2** can react with alkyl halides and amines to form compounds of the invention with structures **11** and **12**. Alcohol **2** can also be transferred into a halide or its equivalents ($X = Cl, Br, I, OMs, \text{ or } OTs$), followed by alkylation with a variety of alkylating reagents to
- 10 afford compounds of the invention with structures **13**, **14**, and **15**.

Scheme 2

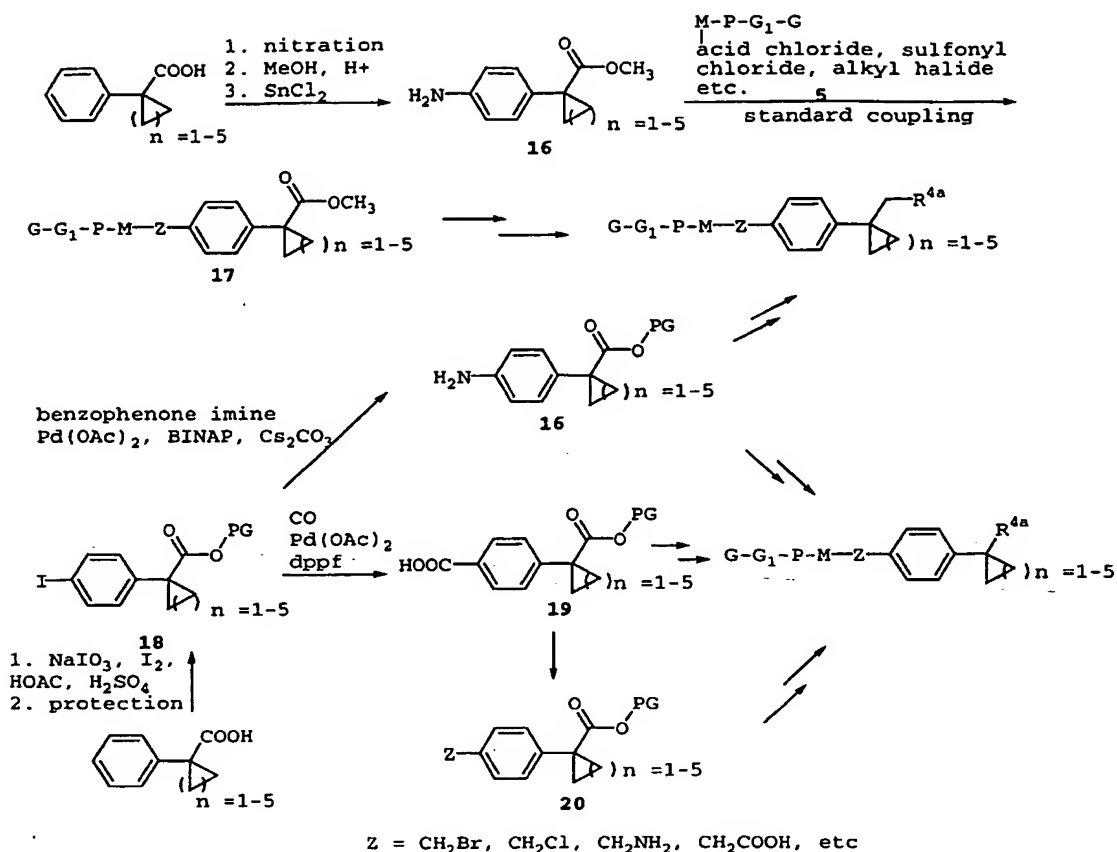


Other compounds of the present invention where Y is a
 5 cycloalkyl derivative can be prepared using commercial
 available 1-phenylcycloalkylcarboxylic acids (or 1-
 phenylcycloalkylcarbonitriles) as the starting material as

illustrated in Scheme 3. Thus, nitration, followed by reduction of the NO₂ group and protection of the acid group can provide the A-B precursor 16, which can be coupled with 5 using standard coupling conditions to provide 17.

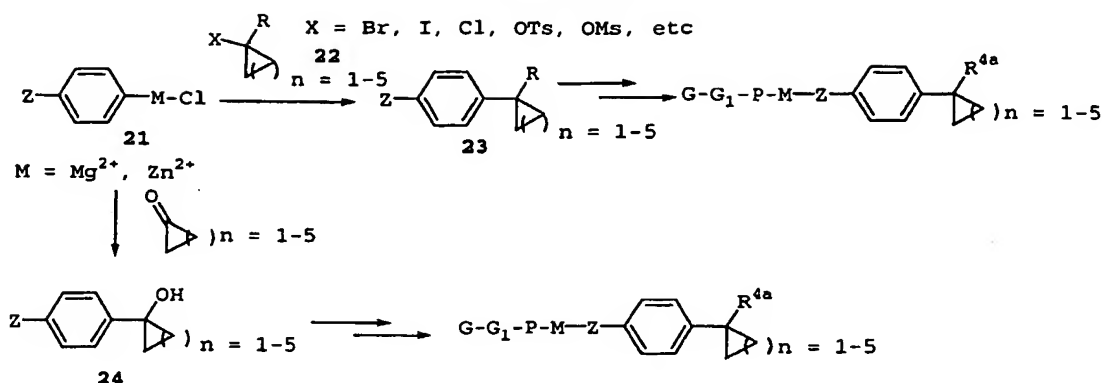
- 5 Alternatively, iodination will provide the desired *para*-substituted compound 18, which can in turn be transformed to the amine 16 via Buchwald palladium-catalyzed amination (*Tetrahedron Lett.* 1997, 38, 6367-6370) and the acid 19 via palladium-catalyzed carboxylation (CO, Pd(OAc)₂, dppf).
- 10 Additional Z-linkers to the A-B intermediates can be synthesized by chemical manipulation of the amino and carboxylic acid functionality in 16 and 19, respectively. Compound 19 can be homologated via the Arndt-Eistert methodology to afford other A-B intermediates in 20.
- 15 Alternatively, the acid functionality in 19 can be reduced to the alcohol that in turn can be converted to a variety of A-B intermediates 20 by procedures known to those skilled in the art. Further elaboration of these intermediates using the methods described above and by
- 20 those skilled in the art should provide compounds of the present invention.

Scheme 3



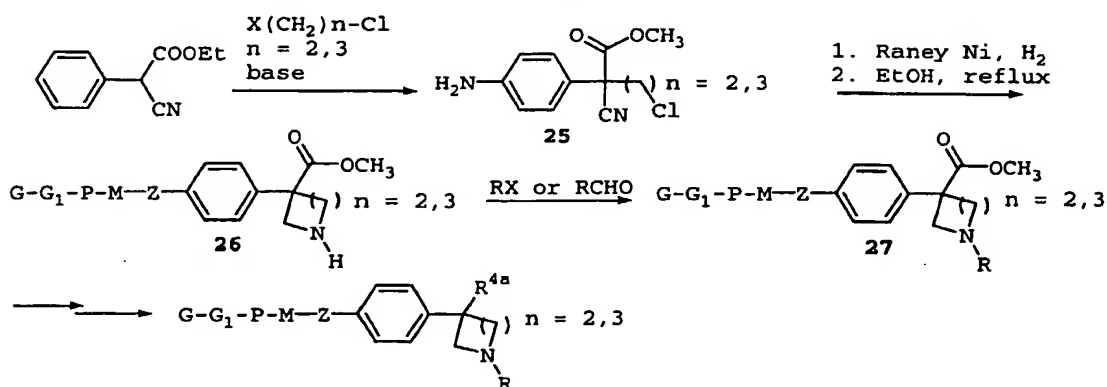
Other compounds of the present invention where Y is a
 5 cycloalkyl derivative can be prepared using organometallic
 reagents (Zn, Mg, etc) **21** as starting materials as shown in
 Scheme 4. Reaction of **21** with properly substituted
 cycloalkyl halides **22** (X = Cl, Br, I, OMs, OTs, etc.) using
 Pd(dba)₂/1,2-bis(diphenylphosphino)ethane (dppe) or
 10 NiCl₂(PPh₃)₂ as catalyst system will provide intermediate
23. Alternatively, Grignard reaction of **21** with cycloalkyl
 ketones will provide intermediate **24**. Further elaboration
 of **23** and **24** using the methods described above and by those
 known in the art should provide compounds of the present
 15 invention.

Scheme 4



Compounds of formula I where Y is a pyrrolidine or piperidine derivative can be prepared as shown in Scheme 5. Thus, phenylcyanoacetate can be alkylated with X-(CH₂)_n-Cl (X, Y = Br, I, OMs, OTs, etc, n = 2,3) to provide the chloronitrile 25, which can be reduced to the corresponding primary amine, followed by cyclization in refluxing EtOH to form 3-pyrrolidine or 3-piperidine derivatives 26. Alkylation or reductive amination can provide the N-substituted intermediate 27. Further elaboration using the methods described above and by those skilled in the art should provide compounds of the present invention.

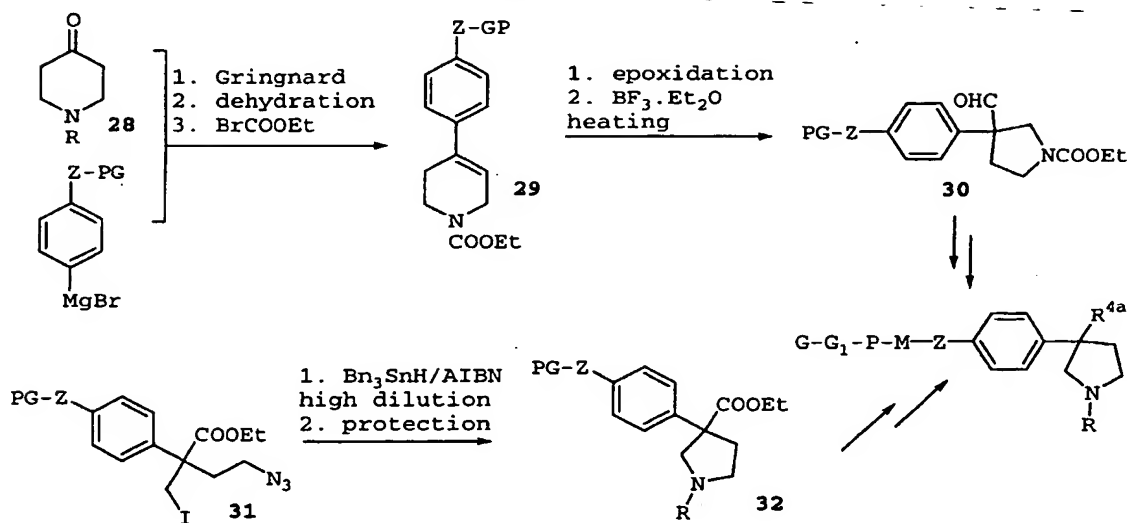
Scheme 5



Compounds of formula I where Y is a pyrrolidine derivative can also be prepared as illustrated in Scheme 6. The Grignard reaction of 1-substituted 4-piperidone 28 with

the appropriate arylmagnesium halide followed by dehydration will give tetrahydropyridine derivative **29**. Epoxidation, followed by rearrangement with heating in boron trifluoride etherate (Chem. Pharm. Bull. 28(5), 1387-1393 (1980)) will provide pyrrolidine aldehyde **30**. Alternatively, radical cyclization of alkyl azide **31** (Tetrahedron Lett. 1997, 38, 3915-1918) can provide the pyrrolidine intermediate **32**. Further elaboration of these intermediates using the methods described above and by those skilled in the art should provide compounds of the present invention.

Scheme 6

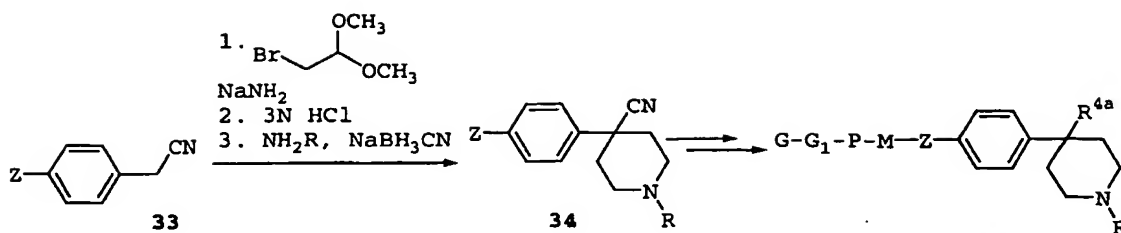


15

Compounds of formula I where Y is a 4-piperidine derivative can be prepared using 2-aryl acetonitriles **33** as starting materials as shown in Scheme 7. Dialkylation of **33** with bromoacetaldehyde dimethyl acetal, followed by hydrolysis of the acetals and reductive amination will give the 4-aryl-4-cyanopiperidine **34**. Further elaboration of these intermediates using the methods described above and by those skilled in the art should provide compounds of the present invention.

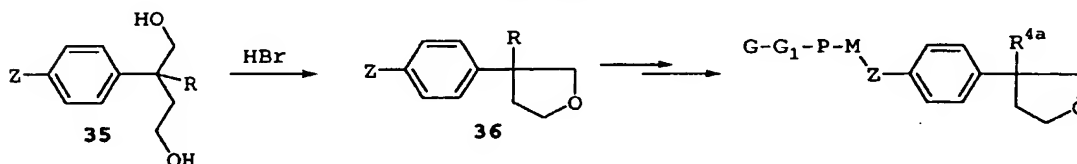
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Scheme 7



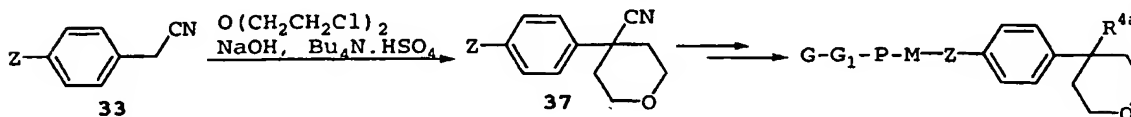
Compounds of formula I where Y is a 4-tetrahydrofuran derivative can be prepared using diol 35 as the starting material as illustrated in Scheme 8. Cyclization of 35 with HBr will give the 4-aryl-4-substituted tetrahydrofuran 36. Further elaboration using the methods described above and by those skilled in the art should provide compounds of the present invention.

Scheme 8



Compounds of formula I where Y is a 4-tetrahydropyran derivative can be prepared using 2-aryl acetonitriles 33 as starting materials as shown in Scheme 9. Alkylation of 33 with di-2-chloroethyl ether will give the 4-aryl-4-cyanotetrahydropyran 37. Further elaboration using the methods described above and by those skilled in the art should provide compounds of the present invention.

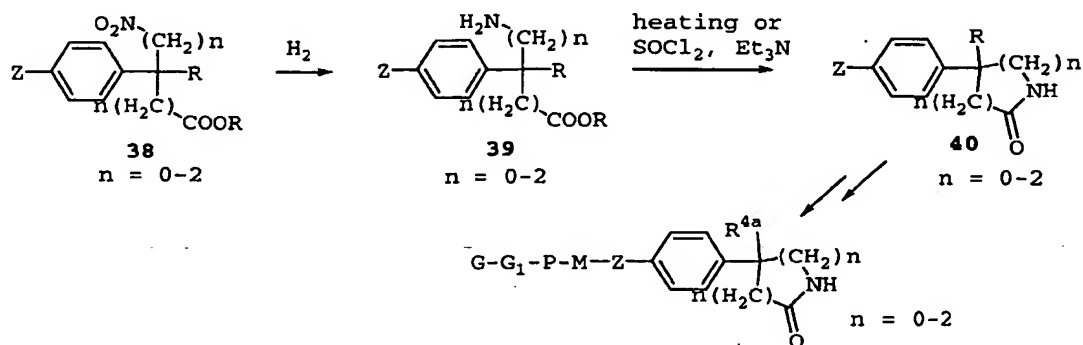
Scheme 9



Compounds of formula I where Y is a lactam derivative can be prepared using intermediate 38 as the starting

material as shown in Scheme 10. Reduction of NO₂ group or nitrile group will provide the primary amine **39**, which can be coupled intramolecularly with the acid or ester to form the lactam **40**. Further elaboration using the methods
 5 described above and by those skilled in the art should provide compounds of the present invention.

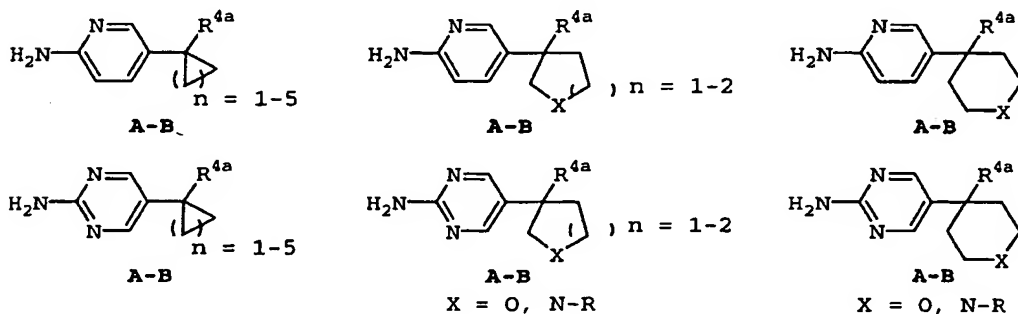
Scheme 10



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Aminopyridyl and aminopyrimidyl A-B analogs (see structures in Scheme 11) can be prepared using routes similar to those of Schemes 2-10 and by those skilled in the art. These intermediates can then be further
 15 manipulated to compounds of this invention with formula I via procedures previously described.

Scheme 11

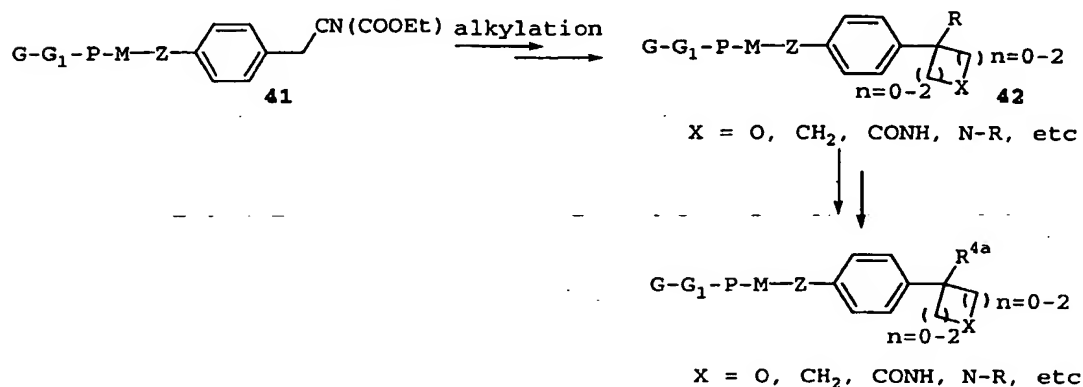


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Compounds of formula I (Scheme 1) where P is fused onto ring M can be prepared as outlined in Schemes 12 and 13, and via standard methods known to those skilled in the

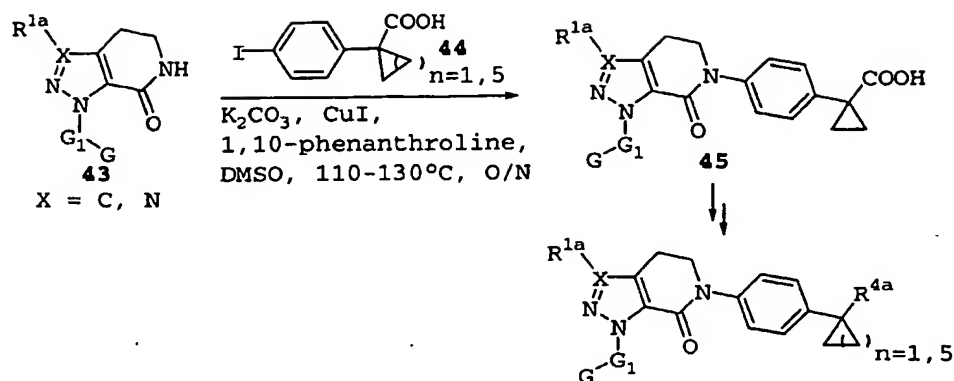
art. The ester or nitrile intermediates **41** illustrated in these Scheme 12 can be subjected to alkylation conditions, followed by other manipulations as described in Schemes 2-10. Further elaboration of intermediates **42** to incorporate the appropriate R^{4a} groups using the methods described above and by those skilled in the art should provide compounds of the present invention.

Scheme 12



Scheme 13 illustrates the synthesis of compounds of formula I (Scheme 1) when P-M moiety **43** is a bicyclic lactam moiety. Thus, the iodo A-B intermediate **44** will
15 react with **43** under Buchwald modified Ullman reaction (*J. Am. Chem. Soc.* **2001**, 123, 7727) using CuI and 1,2-cyclohexyldiamine or 1,10-phenanthroline as the catalyst system to provide **45** in high yields. Further elaboration of **45** to incorporate the appropriate R^{4a} groups art should
20 provide compounds of the present invention by using the methods described above and by those skilled in the art.

Scheme 13



Schemes 2-13 describe how to make the A-B moieties of
 5 the present invention and how to couple them to prepare
 compounds of the present invention. Schemes 2-13 describe
 A-B wherein B is $Y-R^{4a}$ and Y is a cycloalkyl or
 heterocyclyl. Compounds of the present invention wherein Y
 is CY^1Y^2 can be made analogously to the
 10 cycloalkyl/heterocyclyl compounds of Schemes 2-13. For
 example, in Scheme 2, instead of intermediate 1 being a
 cycloalkyl intermediate, it can be Y^1Y^2 disubstituted
 intermediate. This intermediate could be made by a number
 of methods including di-substituting the starting 4-
 15 nitrophenylacetonitrile by reaction with a base and a Y^1 -
 leaving group and a Y^2 -leaving group. One of ordinary
 skill in the art would recognize that other routes to the
 Y^1Y^2 disubstituted intermediate are available. The
 remainder of the chemistry shown in Scheme 3 will then
 20 follow.

In Scheme 3, instead of use the starting 1-
 phenylcycloalkylcarboxylic acids or 1-
 phenylcycloalkylcarbonitriles of Scheme 3, one could use
 the corresponding Y^1Y^2 disubstituted intermediates. Just
 25 like in Scheme 2, these intermediates could be prepared by
 di-substituting a phenylcarboxylic acid or
 phenylcarbonitrile. One of ordinary skill in the art would
 recognize that other routes to these types of Y^1Y^2

disubstituted intermediate are also available. The remainder of the chemistry shown in Scheme 3 will then follow.

The compounds of this invention and the intermediates described above wherein the B group contains an oxidizable group can be oxidized, e.g., N to N-oxide.

In the above Schemes, the Z group may or may not be present depending on how the A-B group is coupled. The coupling portion of the A-B group could (a) be displaced by the incoming Z or M group, (b) become the Z group, or (c) be incorporated into ring M.

The remaining portions of the compounds of the present invention, G-G₁-P-M-Z, G-G₁-M-P-Z, G-G₁-P-M, G-G₁-M-P, G-G₁-M-Z, and G-G₁-M, can be prepared using methods known to those of ordinary skill in the art. All of the following patents and publications are incorporated herein by reference. For compounds wherein ring P is absent and ring M is a 5-, 6-, or 7-membered ring, one of ordinary skill in the art can look to US 5,939,418, US 5,925,635, US 6,057,342, US 6,187,797, US 6,020,357, US 6,060,491, US 5,998,424, US 6,191,159, WO98/57951, WO99/32454, WO00/039108, WO00/059902, WO01/32628, WO01/005785, WO02/00651, WO02/102380, and WO02/00647 for starting materials and intermediates to which the present B and/or A-B groups can be coupled. For compounds wherein ring P is fused to ring M (i.e., a bicyclic moiety is present), one of ordinary skill in the art can look to WO00/39131, WO02/094197, USSN 10/104,467, USSN 10/105,477, and WO02/00655 for starting materials and intermediates to which the present B and/or A-B groups can be coupled.

For compounds wherein G is a ring substituted with a basic moiety, one of ordinary skill in the art can look to US 5,939,418, US 5,925,635, US 6,057,342, US 6,187,797, US 6,020,357, US 6,060,491, US 6,191,159, WO98/57951, WO99/32454, WO00/059902, WO01/32628, WO00/39131, WO02/00651, WO02/102380, WO02/094197, USSN 10/104,467, and USSN

10/105,477 for starting materials and intermediates to form the present G-G₁-P-M-Z, G-G₁-M-P-Z, G-G₁-P-M-Z-A, and/or G-G₁-M-P-Z-A groups to which the present B and/or A-B groups can be coupled. For compounds wherein G is a ring

5 substituted with a non-basic group, one of ordinary skill in the art can look to US 5,998,424, WO00/39131, WO00/059902, WO01/32628, WO02/00651, WO02/102380, WO02/094197, USSN 10/104,467, and USSN 10/105,477 for starting materials and intermediates to form the present G-

10 G₁-P-M-Z, G-G₁-M-P-Z, G-G₁-P-M-Z-A, and/or G-G₁-M-P-Z-A groups to which the present B and/or A-B groups can be coupled. For compounds wherein G is a bicyclic moiety, one of ordinary skill in the art can look to WO98/57951

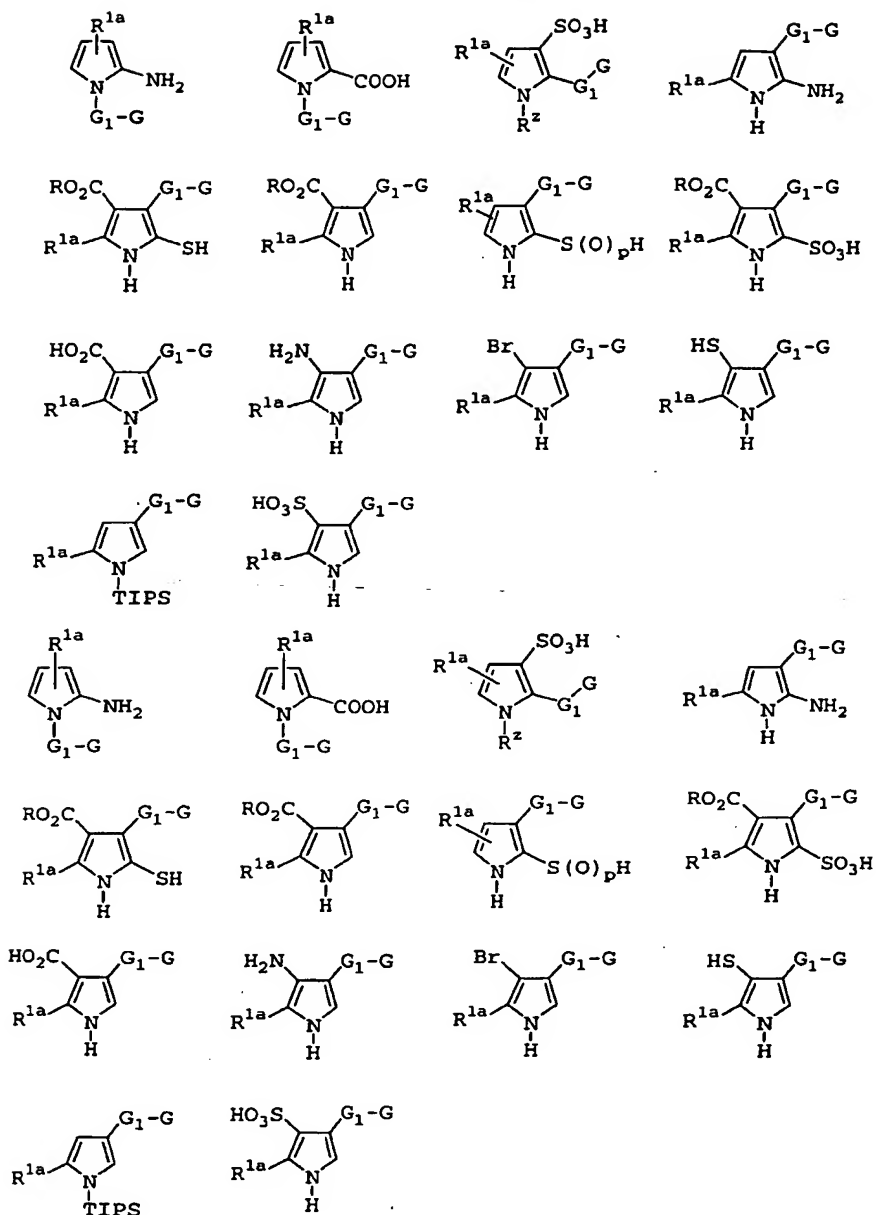
15 WO00/039108, WO00/39131, WO02/00651, WO02/102380, WO02/094197, USSN 10/104,467, and USSN 10/105,477 for starting materials and intermediates to form the present G-G₁-P-M-Z, G-G₁-M-P-Z, G-G₁-P-M-Z-A, and/or G-G₁-M-P-Z-A groups to which the present B and/or A-B groups can be coupled. For compounds wherein A is an indoline or similar

20 bicycle, one of ordinary skill in the art can look to WO01/005785 for starting materials and intermediates to which the present B group can be coupled or from which the present A-B groups can be formed. Scheme 14 illustrates some of the numerous pyrrole intermediates that can be used

25 to prepare compounds of the present invention (R^z is the point of attachment for Z-A-B and can be H, a protecting group, a group modifiable to Z or Z-A, Z, Z-A, or A). These intermediates are described in the above-noted patents and publications.

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Scheme 14

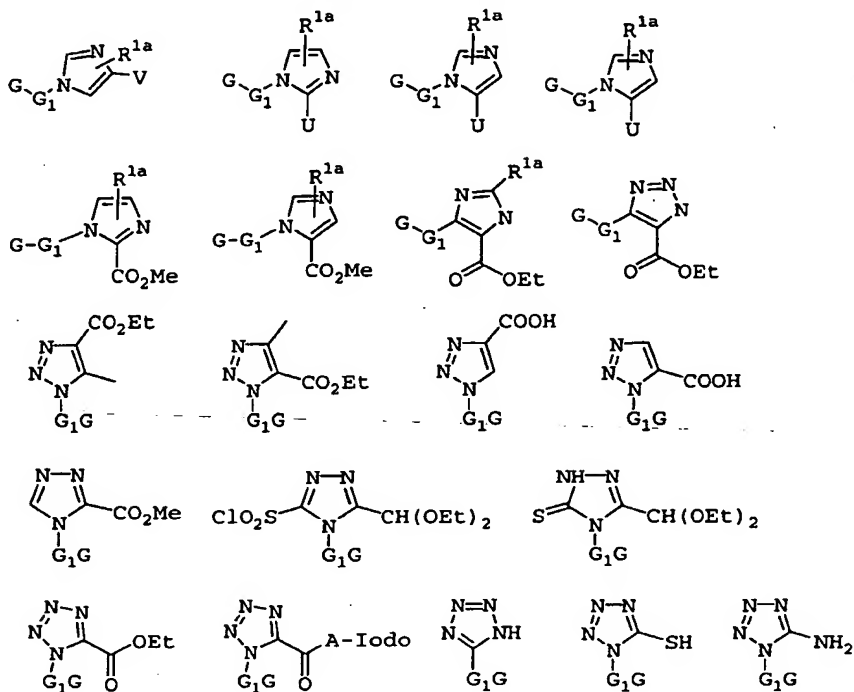


5 Scheme 15 illustrates some of the numerous imidazole, triazole, and tetrazole intermediates that can be used to prepare compounds of the present invention. These intermediates are described in the above-noted patents and publications. In Scheme 15, V is nitro, amino, thio, hydroxy, sulfonic acid, sulfonic ester, sulfonyl chloride, ester, acid, or halide. In Scheme 15, U is aldehyde,

10

ester, acid, amide, amino, thiol, hydroxy, sulfonic acid, sulfonic ester, sulfonyl chloride, or methylene halide.

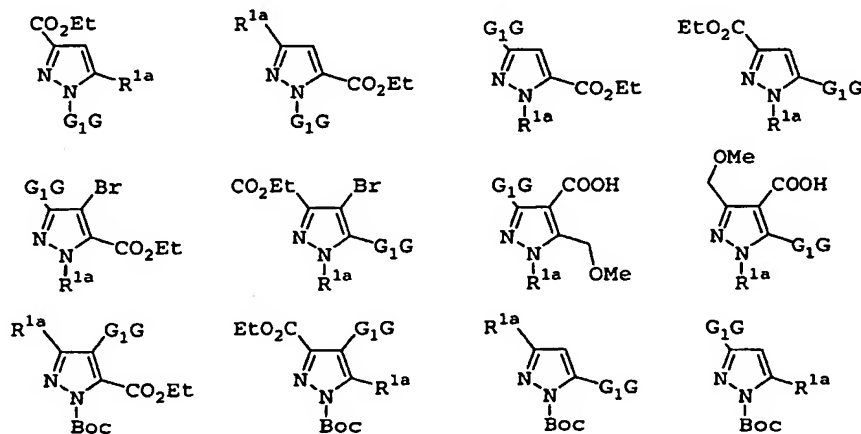
Scheme 15



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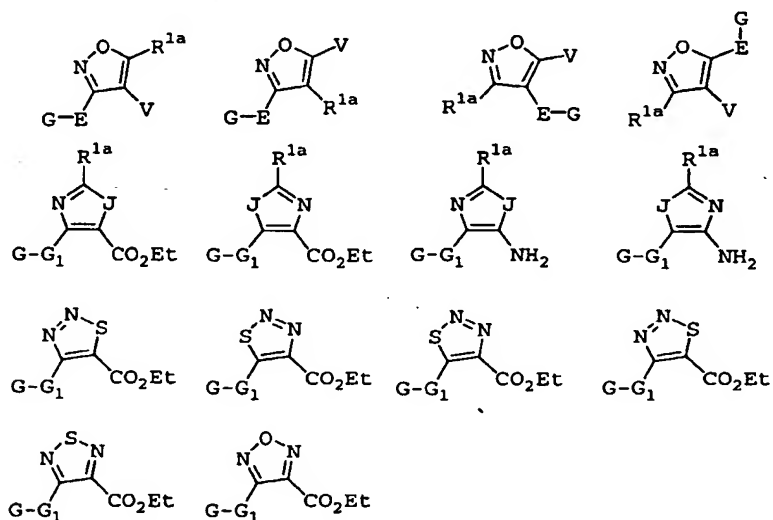
Scheme 16 shows some of the numerous pyrazole intermediates that can be used to prepare compounds of the present invention. These intermediates are described in the above-noted patents and publications.

Scheme 16



Scheme 17 depicts some of the numerous oxazole, thiazole, isoxazole, oxadiazole, and thiadiazole intermediates that can be used to prepare compounds of the present invention. These intermediates are described in the above-noted patents and publications. In Scheme 17, V is nitro, amino, ester, or acid.

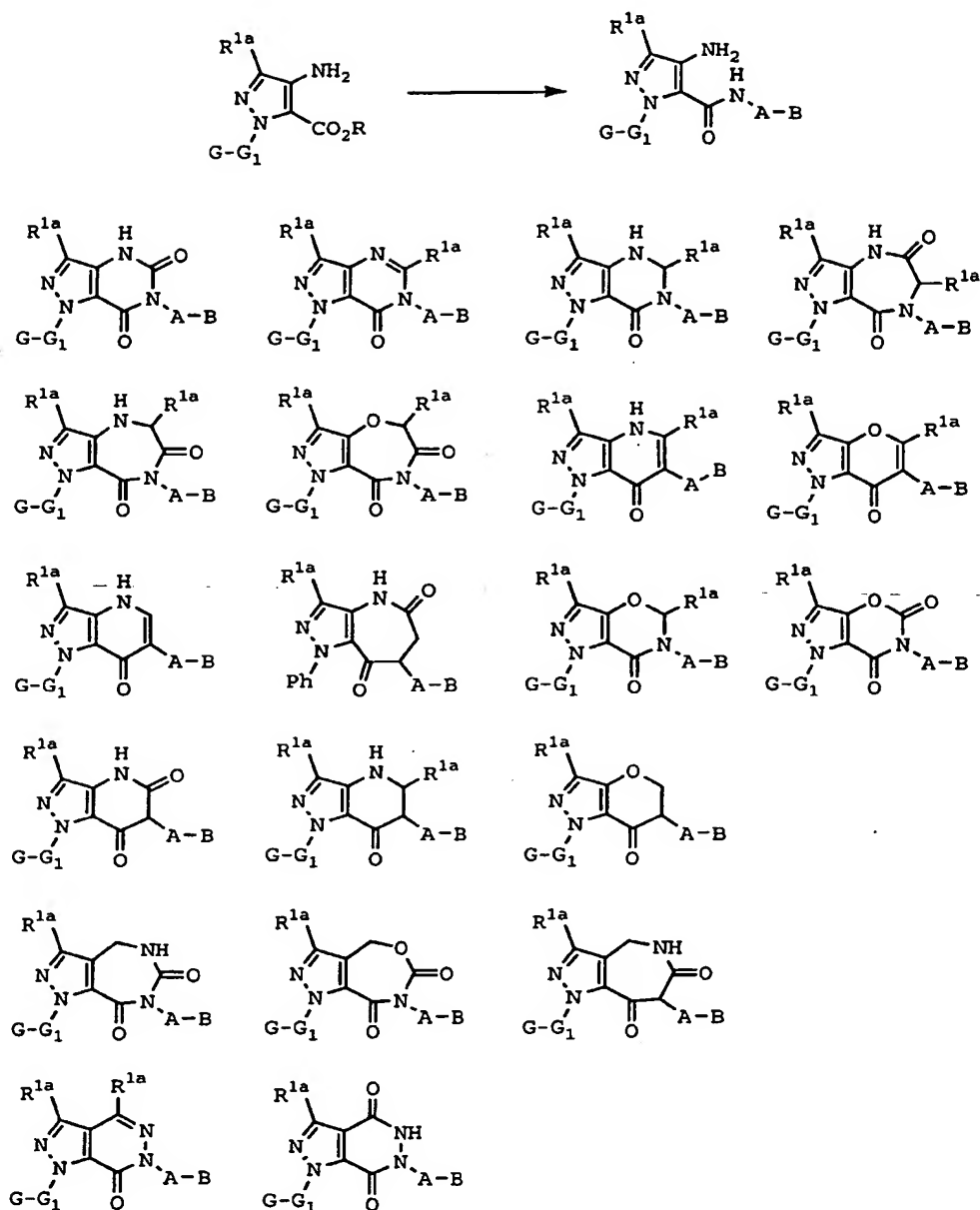
Scheme 17



10

Scheme 18 illustrates two intermediates useful for making a compound of the present invention wherein ring P is fused to ring M. Scheme 18 also illustrates a number of bicyclic compounds that can be made from these intermediates or derivatives thereof. These intermediates and their modification are described in the above-noted patents and publications.

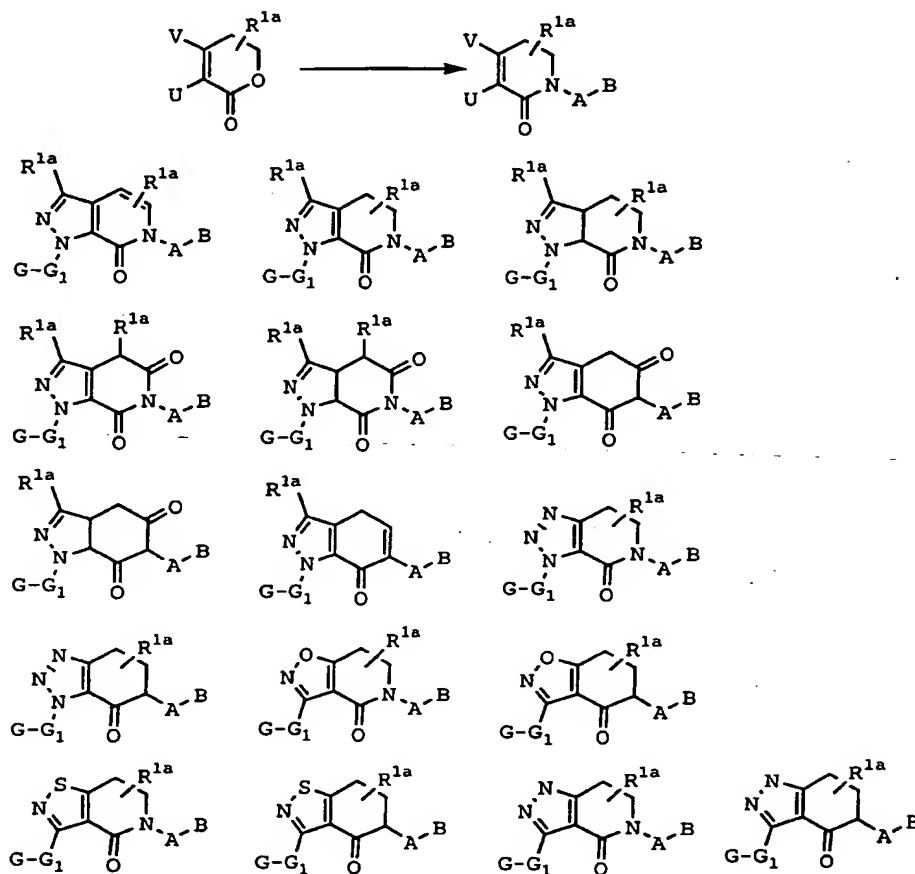
Scheme 18



Scheme 19 depicts another intermediate useful for making a compound of the present invention wherein ring P is fused to ring M. Scheme 19 also illustrates a number of bicyclic compounds that can be made from this intermediate or derivatives thereof (e.g., the corresponding cyclohexenone). In Scheme 19, U is OH or morpholine and V is H or $C(O)R^{1a}$. This intermediate, derivatives thereof,

and their modification are described in the above-noted patents and publications.

Scheme 19

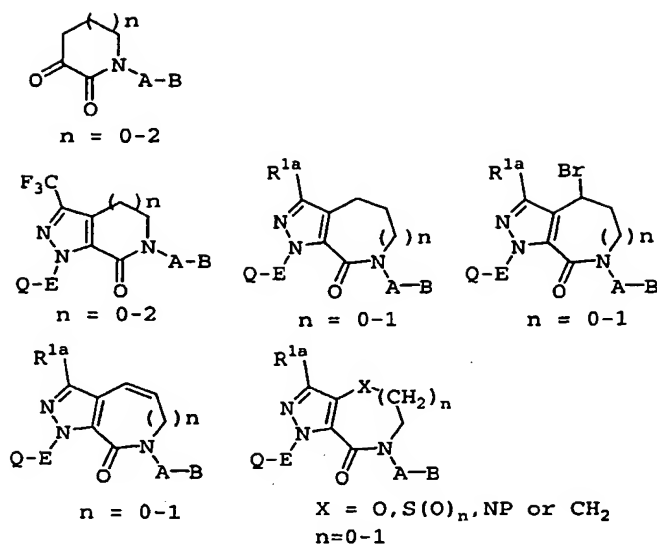


5

Scheme 20 shows another intermediate useful for making a compound of the present invention wherein ring P is fused to ring M. Scheme 20 also illustrates a number of bicyclic compounds that can be made from this intermediate or derivatives thereof. This intermediate, derivatives thereof, and their modification are described in the above-noted patents and publications.

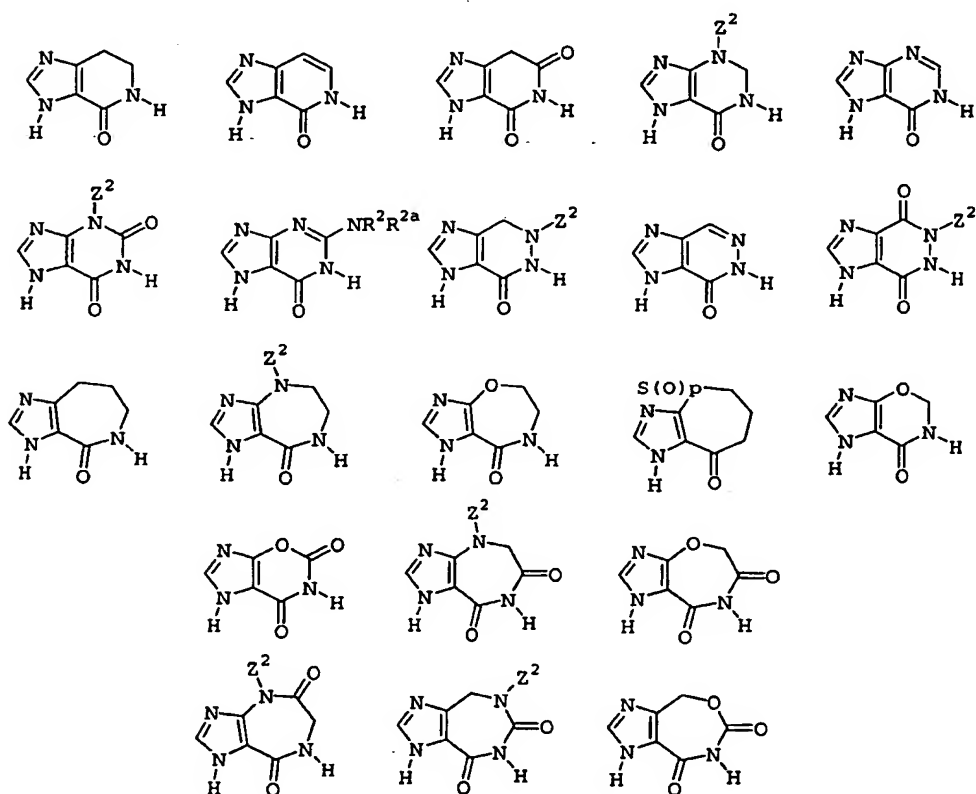
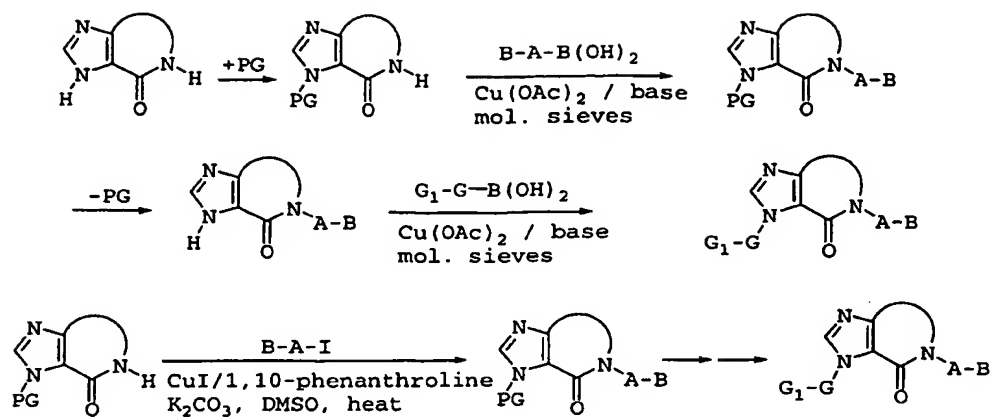
10

Scheme 20



Scheme 21 illustrates a number of other bicyclic rings
 5 that are considered to be part of the present bicyclic
 group, rings P-M. Scheme 21 also describes a method of
 converting the shown rings to compounds of the present
 invention. As one of ordinary skill in the art would
 recognize, this method would be applicable to other
 10 heterobicyclics not shown.

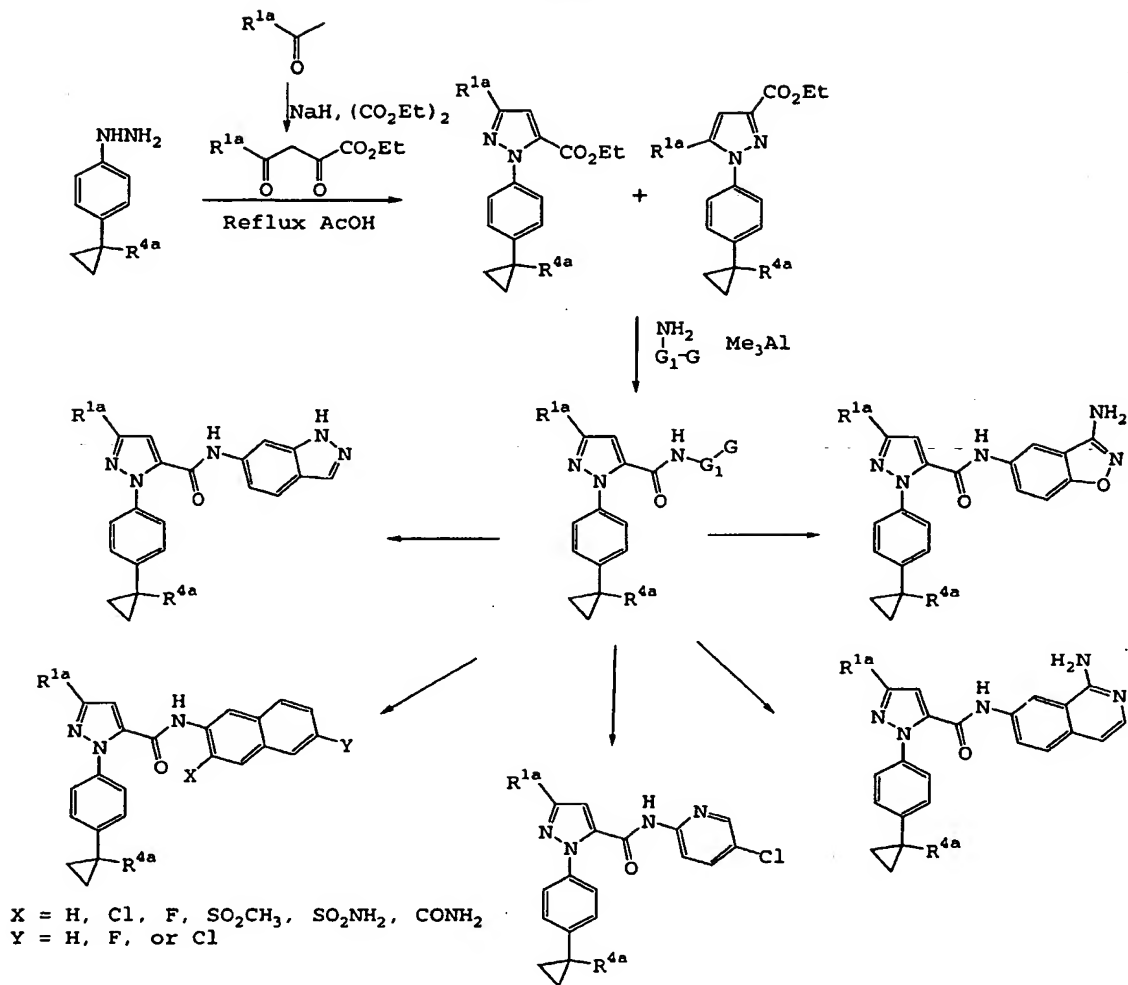
Scheme 21



Other useful pyrazole intermediates wherein G₁ is an amide are exemplified in Scheme 22. Compounds of the present invention wherein the G₁ group is other than an amide can be easily manipulated to other linker functionalities according to the methodologies known in the art, including the methodologies outlined in WO98/28269 and

WO98/28282, the contents of both are incorporated herein by reference.

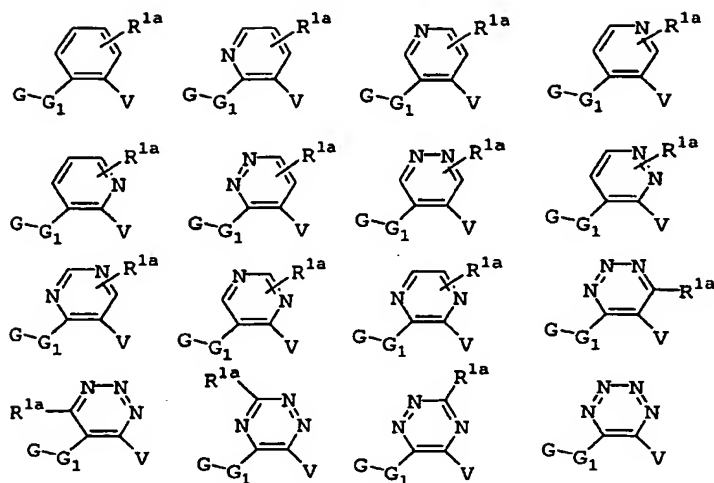
Scheme 22



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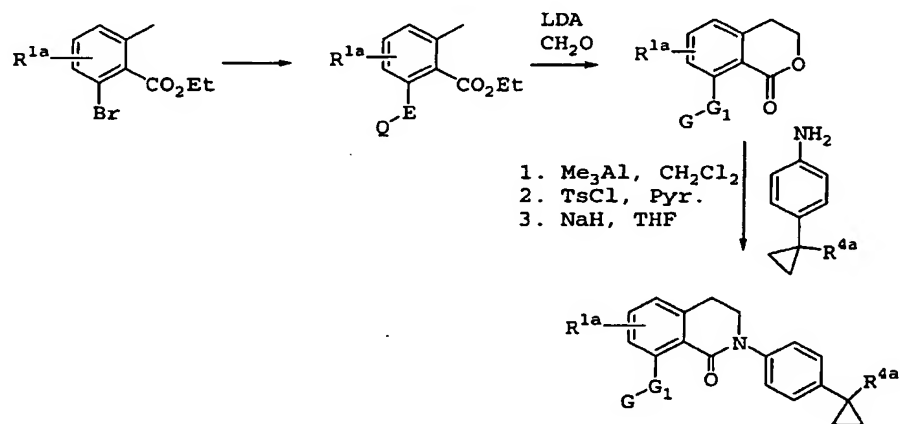
Scheme 23 depicts some of the numerous 6-membered aromatic ring intermediates that can be used to prepare compounds of the present invention. These intermediates are described in the above-noted patents and publications. In Scheme 23, V is nitro, protected sulfonamide, or ester group and is a precursor of group Z of the present invention.

Scheme 23



Benzo fused dihydro-pyridone intermediates of the present invention can be prepared from readily available starting materials as shown in Scheme 24.

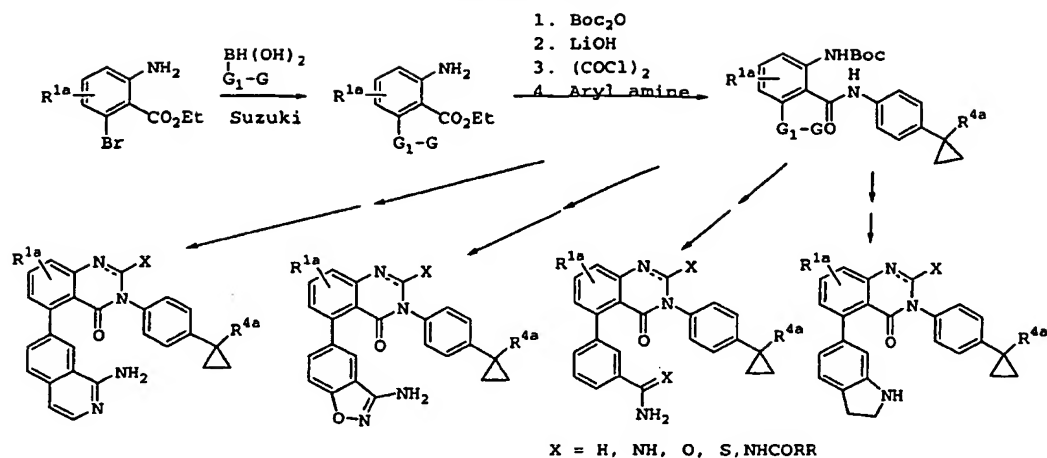
Scheme 24



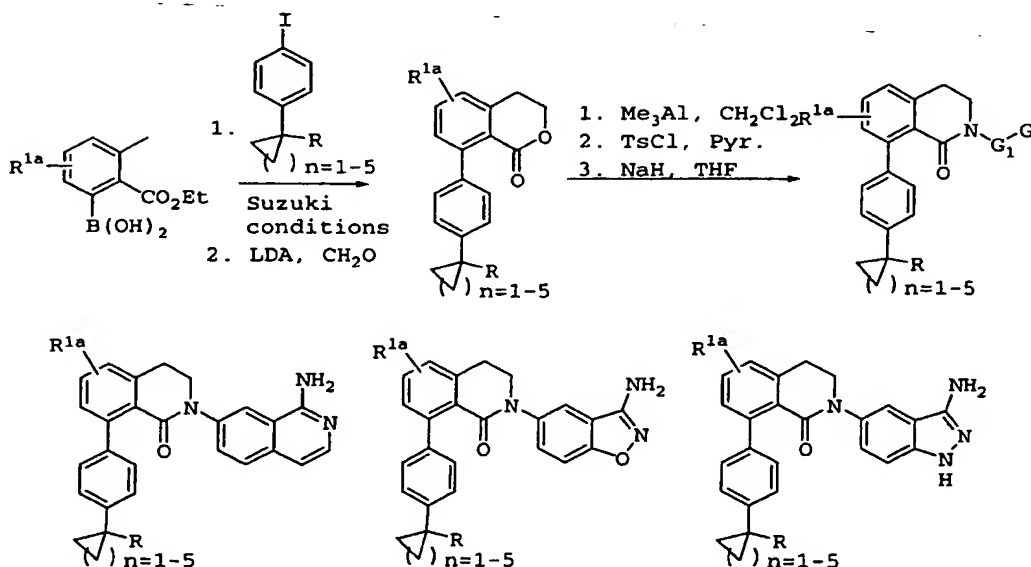
10

Other benzo-bicyclics can also be obtained as shown in Schemes 25 and 26.

Scheme 25



Scheme 26

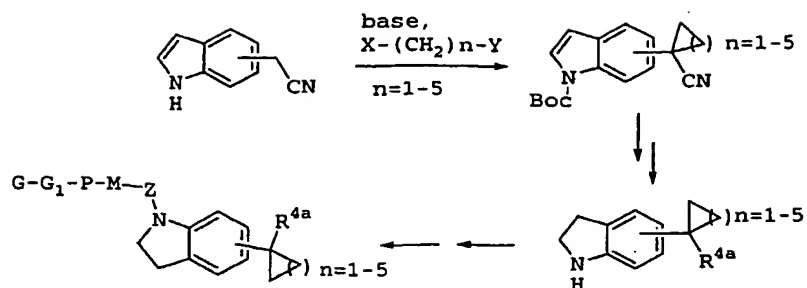


5

Intermediates A-B of the present invention wherein A is indoline can be prepared as shown in scheme 27. This type of intermediate can then be attached to the remainder of the desired compound as described previously.

Alternatively, the indoline can be attached to the other half of the desired compound prior to formation of the carbocyclic or heterocyclic ring.

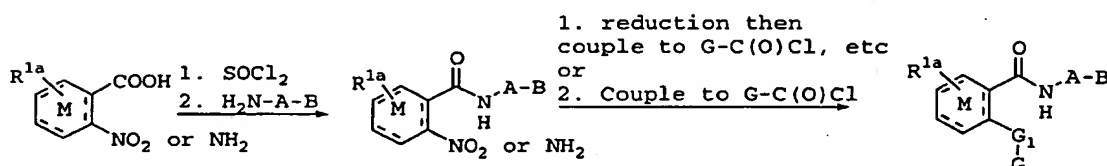
Scheme 27



Compounds of the present invention wherein ring P is
 5 absent and ring M is a six-membered ring can be obtained as
 shown in scheme 28. These types of compounds can be
 obtained from commercially available anthranilic acids or
 their anthranilates. Anthranilic acids or their nitro
 precursors can be coupled with a suitable B-A-NH₂ in
 10 presence of a base such as triethyl amine, pyridine, or
 DMAP. Subsequent coupling with an appropriate acid
 chloride or aniline or aminopyridyl should afford compounds
 of the present invention.

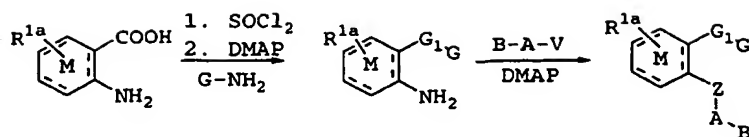
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Scheme 28



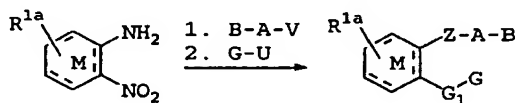
In an analogous fashion the anthranilates can be
 20 coupled with a suitable amine, aniline, or aminopyrimidyl
 to afford the corresponding benzamide. The benzamides can
 then be coupled with an appropriate B-A-V (wherein V is a
 acid chloride derivative, an alkyl halide, or a sulfonyl
 chloride) to afford additional compounds of the present
 25 invention (see scheme 29).

Scheme 29



5 Commercially available ring M derivatives bearing a
 nitro and amino functionality can also be derivatized as
 shown above to afford bisamide analogs. In this case,
 coupling of the aniline with B-A-V (wherein V is an acid
 chloride, a sulfonyl chloride, or an alkylhalide) affords
 10 an intermediate that can be subjected to treatment with an
 appropriate G-U (wherein U is either an acid chloride or an
 alkyl halide) in presence of a suitable base such as DMAP.
 It should be noted that the order of addition of B-A-V and
 G-U can be reversed to obtain other compounds of the
 15 present invention (see scheme 30).

Scheme 30

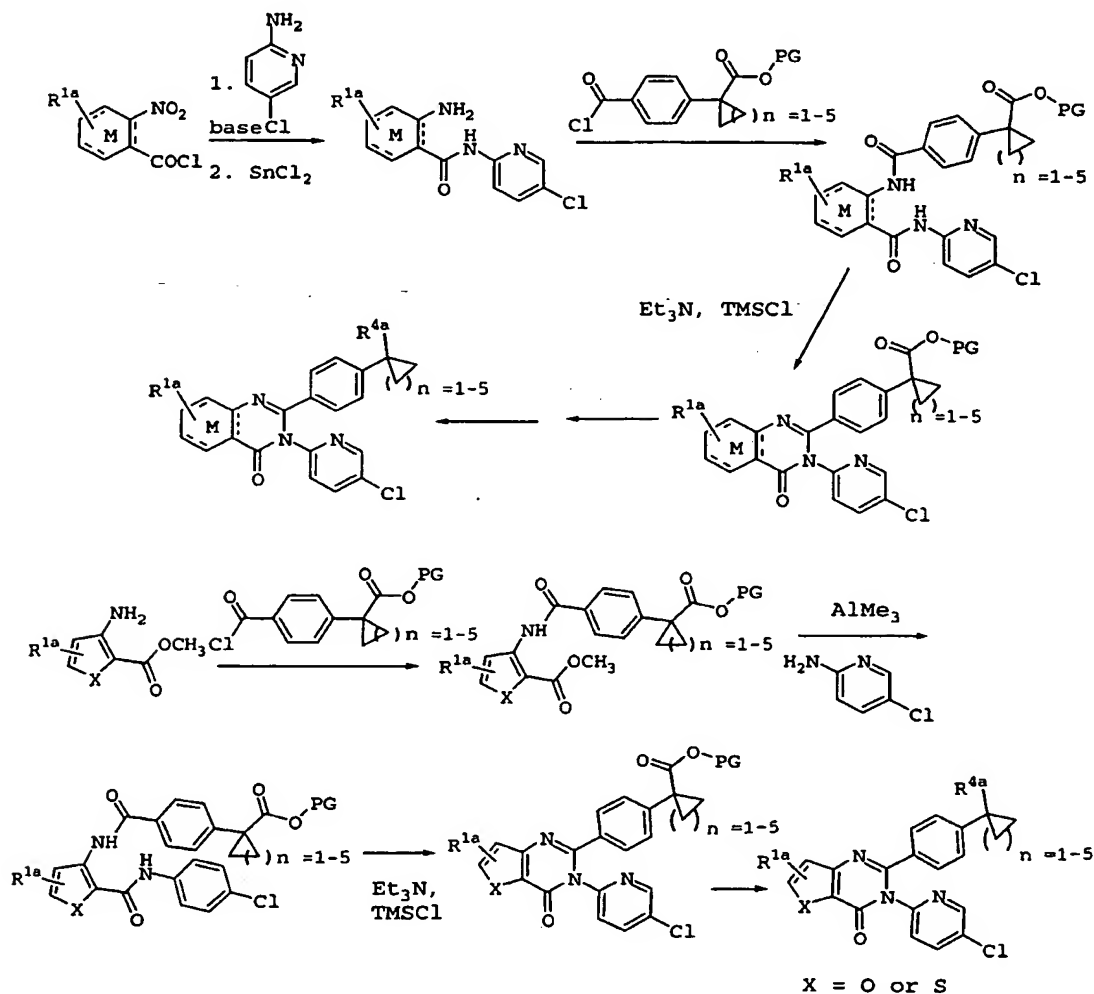


20

It should be noted that the syntheses shown above
 could be modified to use coupling intermediates such as
 Iodo-A-V, wherein V is an acid chloride, amino,
 alkylhalide, or sulfonyl chloride. These in turn could be
 25 coupled to a G-U group. The iodo intermediate could then
 be subjected to Ullman or Buchwald coupling as described
 previously to afford compounds of the present invention.
 The iodo intermediate could also be converted to an amine
 via standard Buchwald conditions to afford the
 30 corresponding anilino intermediate. This in turn could be
 coupled as previously described to afford compounds of the
 present invention.

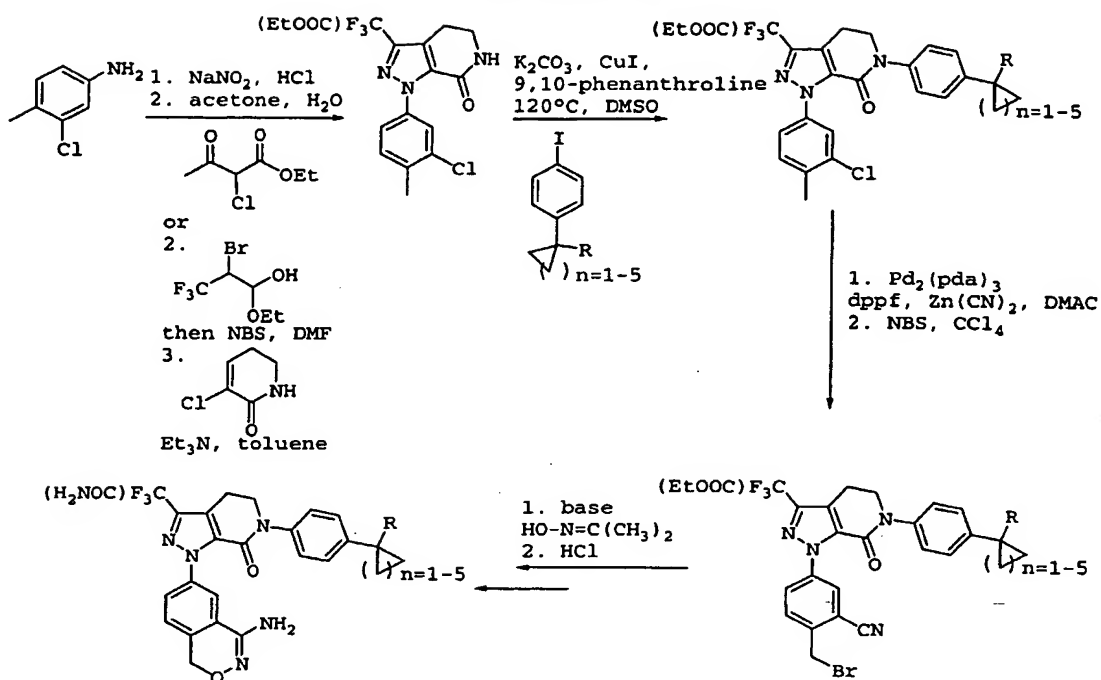
The syntheses of bisamide compounds shown in Schemes 28-30 can also be applied to the syntheses of compounds with ring M as a 5-membered heterocycle. The bisamides can also be further converted into bicyclic pyrimidin-4-ones under acidic conditions as shown in Scheme 31.

Scheme 31



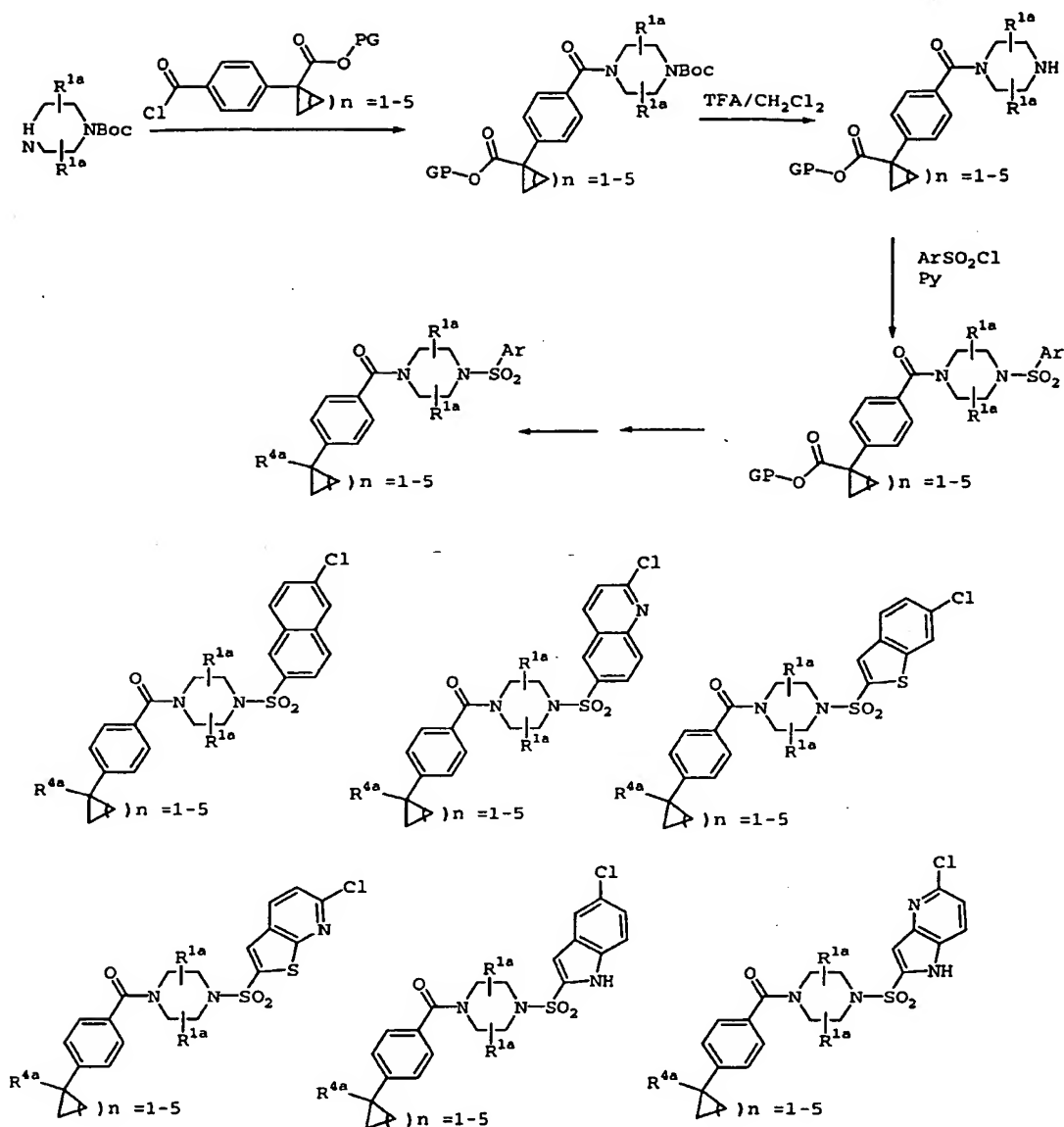
10 Scheme 32 depicts the synthesis of aminobenzines by using the methods described above and by those skilled in the art.

Scheme 32



Scheme 33 illustrates the synthesis of piperidine derivatives by using the methods described above and known by those skilled in the art.

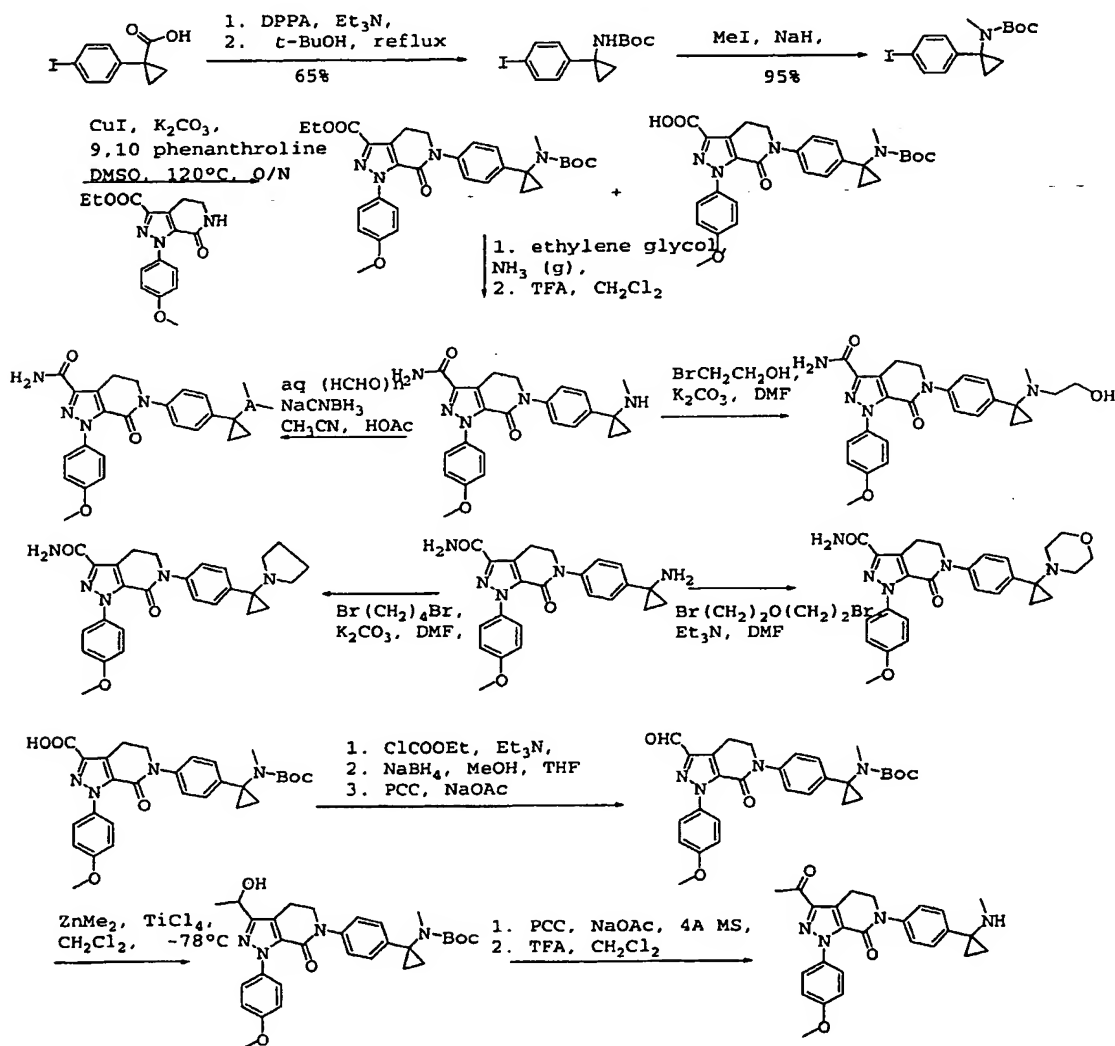
Scheme 33



Scheme 34 depicts the syntheses of phenylcyclopropyl amine derivatives. Starting from 4-iodophenylcyclopropyl carboxylic acid, Curtius rearrangement with DPPA in CH₂Cl₂ at RT followed by heating in *t*-BuOH afforded Boc-protected cyclopropylamine intermediate. This intermediate underwent a sequence of methylation (MeI, NaH, THF), Buchwald Ullman coupling (CuI, K₂CO₃, 9,10-phenanthroline, DMSO), and then amination of the ethyl ester (NH₃ in ethylene glycol) to yield the desired product. Reductive amination with aqueous

formaldehyde and NaBH_3CN in CH_3CN afforded the dimethyl compound. On the other hand, alkylation with bromoethanol, dibromobutane, or dibromoethylether using K_2CO_3 as the base yielded tertiary or cyclic amines, respectively. The C3-methylketone analogue was synthesized through a sequence involving a nucleophilic reaction of ZnMe_2 with the aldehyde in the presence of TiCl_4 .

Scheme 34



10

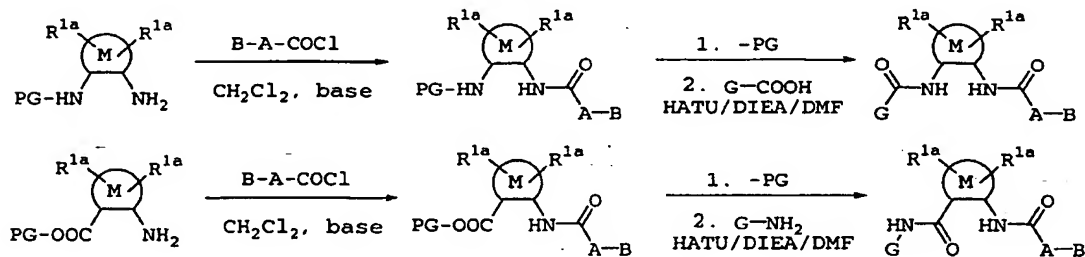
Compounds of the present invention wherein ring P is absent and ring M is a 3-10 membered non-aromatic carbocycle or heterocycle can also be prepared by using the

methods described previously and known to those skilled in the art. Scheme 34 illustrates a number of nonaromatic M rings that are considered to be part of the present invention. Scheme 34 also describes general methods of

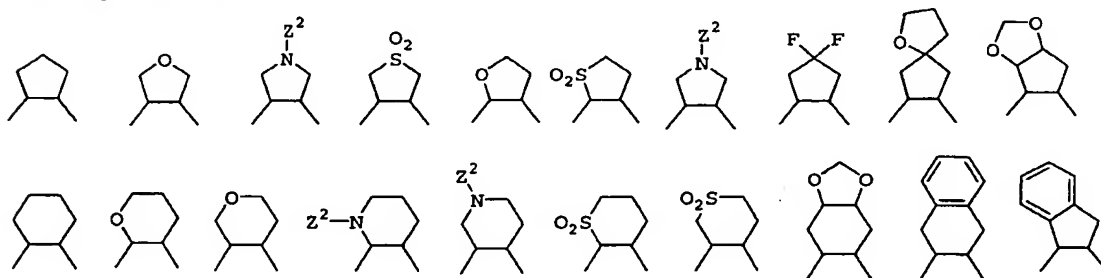
5 converting the shown rings to compounds of the present invention. As one of ordinary skill in the art would recognize, this method would be applicable to other non-aromatic rings not shown.

10

Scheme 35



M rings can be:



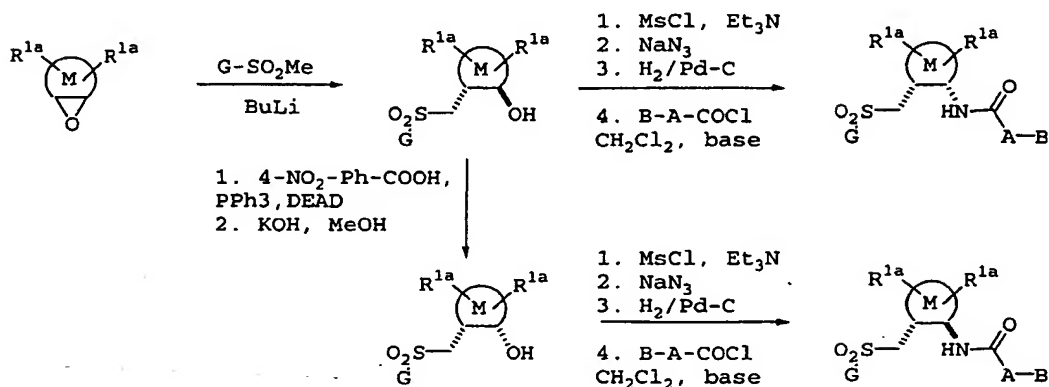
The properly protected, enantiomerically pure cyclic amino acid cores can be obtained via Davies' protocol (*J. Chem. Soc. Perkin Trans I*, **1994**, 1411) or via the reduction of enamines described by Cimarelli, C. et al (*J. Org. Chem.* **1996**, 61, 5557). The corresponding diamino compounds can be obtained via saponification of the ester of the cyclic amino acids followed by Curtius rearrangement. On the other hand, the cyclic diamines can be prepared via

20 literature methods. (See, for example, Skarzewski, J. and Gupta, A. *Tetrahedron: Asymmetry*, **1997**, 8, 1861 and Kim, B. M.; Bae, S. J.; and Seomoon, G., *Tetrahedron Lett.* **1998**, 39, 6921).

A series of compounds of formula I wherein G_1 is 1,1-dioxo-sulfonylmethyl group are prepared following the sequence outlined in Scheme 36.

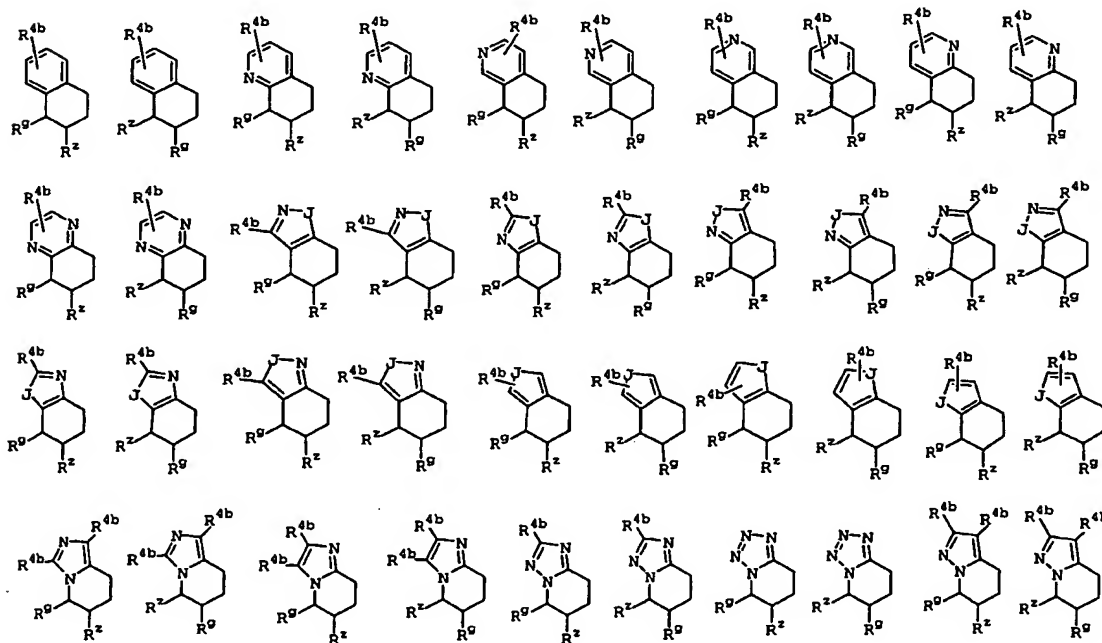
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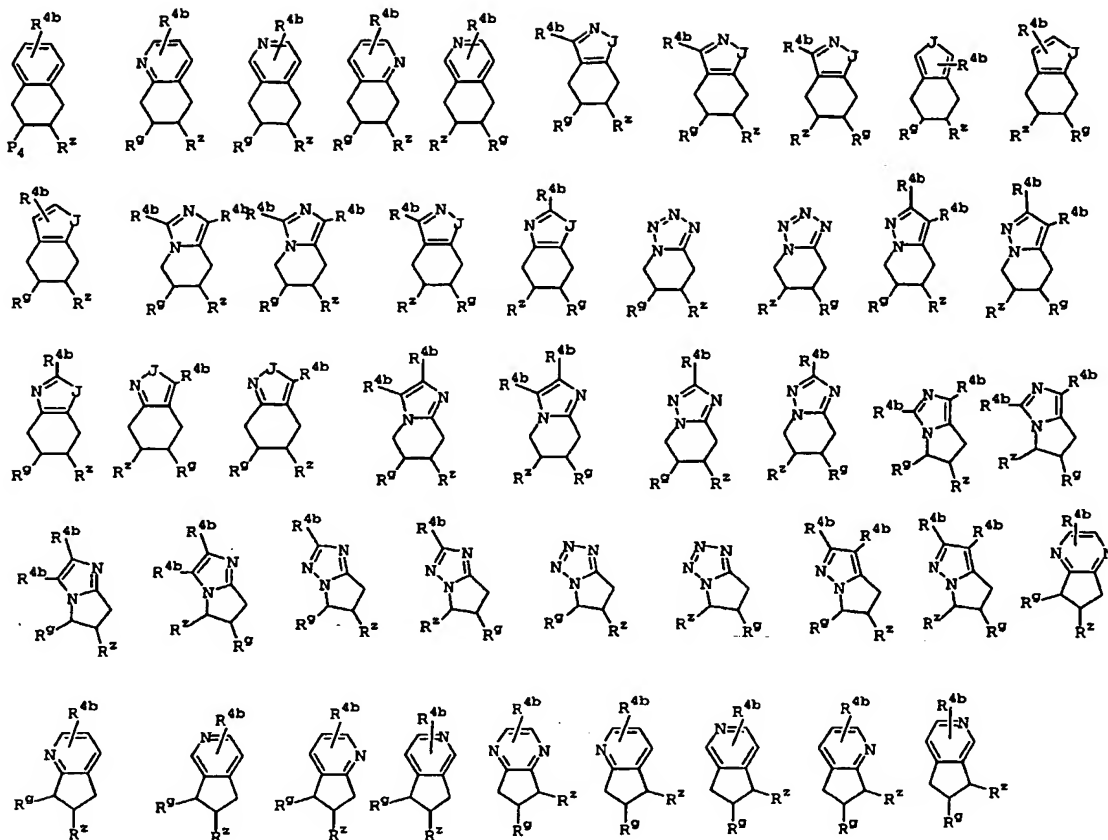
Scheme 36



Scheme 37 illustrates numerous bicyclic M intermediates that can be used to prepare compounds of the present invention. These intermediates can be prepared using methods known to those of ordinary skill in the art and using similar methods described previously.

Scheme 37



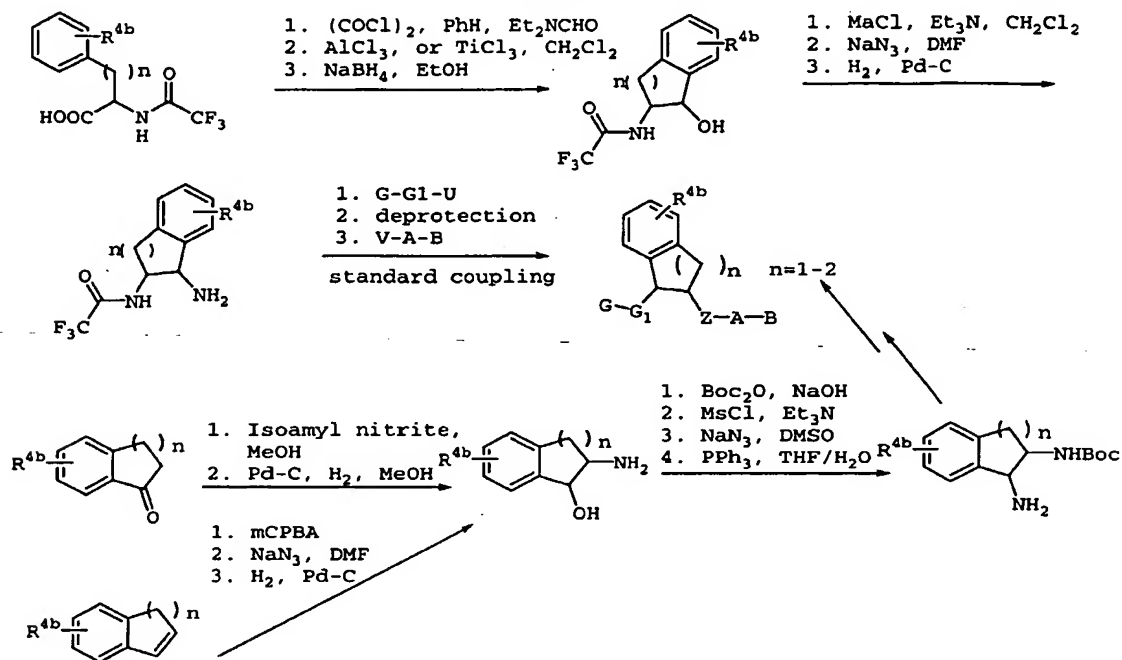


Scheme 38 illustrates the synthesis of benzofused M intermediates of the present invention. The α - or β -amino acid derivatives can undergo Friedel-Crafts reaction followed by reduction to afford the fused ring intermediates. Replacement of the OH group with NH_2 group as described previously, followed by standard coupling reactions will provide the compounds of the present invention. On the other hand, oxime formation of the ketone intermediate followed by reduction with NaBH_4 can provide the amino alcohol intermediate, which can also be obtained via epoxidation of the olefin and then nucleophilic displacement. Protection of the amino group followed by azide displacement of the mesylate and then reduction of the azide group will give the Boc protected diamines. Functional groups U and V can be acid chloride, carboxylic acid, sulfonyl chloride, etc. in formula U-G₁-G

and V-A-B. The compounds of the present invention can be obtained from the mono-Boc protected diamines using methods known to those of ordinary skill in the art and using similar methods described previously.

5

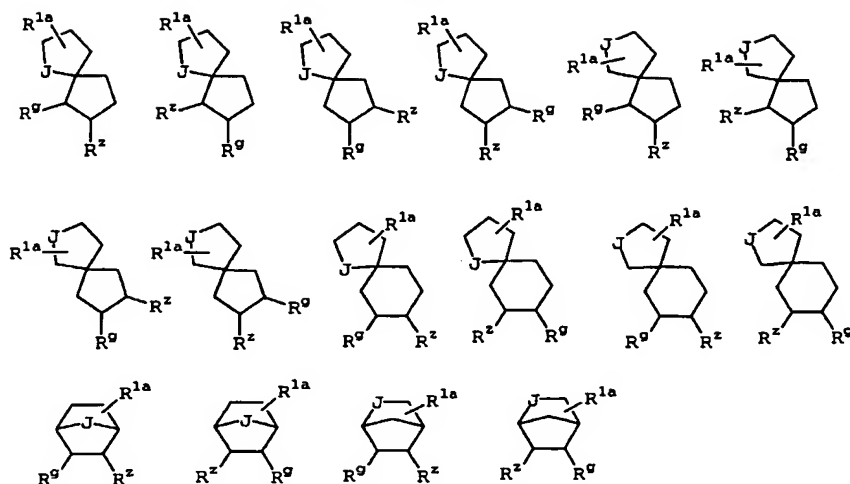
Scheme 38



Scheme 39 depicts numerous spiro and bridged M intermediates that can be used to prepare compounds of the present invention. These intermediates can be prepared using methods known to those of ordinary skill in the art and using the methods described previously.

10

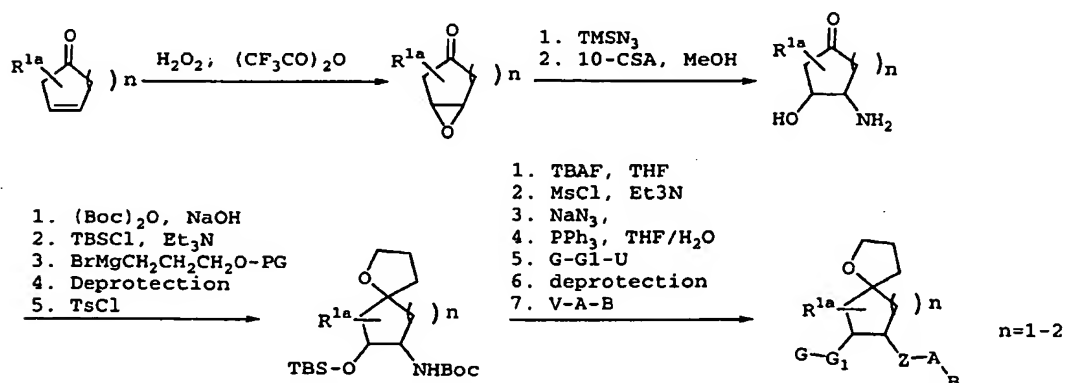
Scheme 39



Scheme 40 depicts the synthesis of spiro intermediates of the present invention. Epoxidation of the olefin followed by displacement with TMSN_3 and reduction with 10-CSA can provide the amino alcohol intermediate. Protection of the amino and alcohol groups followed by nucleophilic addition to the carbonyl group and spiro ring formation can afford the spiro tetrahydrofuran intermediate. This intermediate can undergo a similar sequence of reactions described previously to give compounds of the present invention.

15

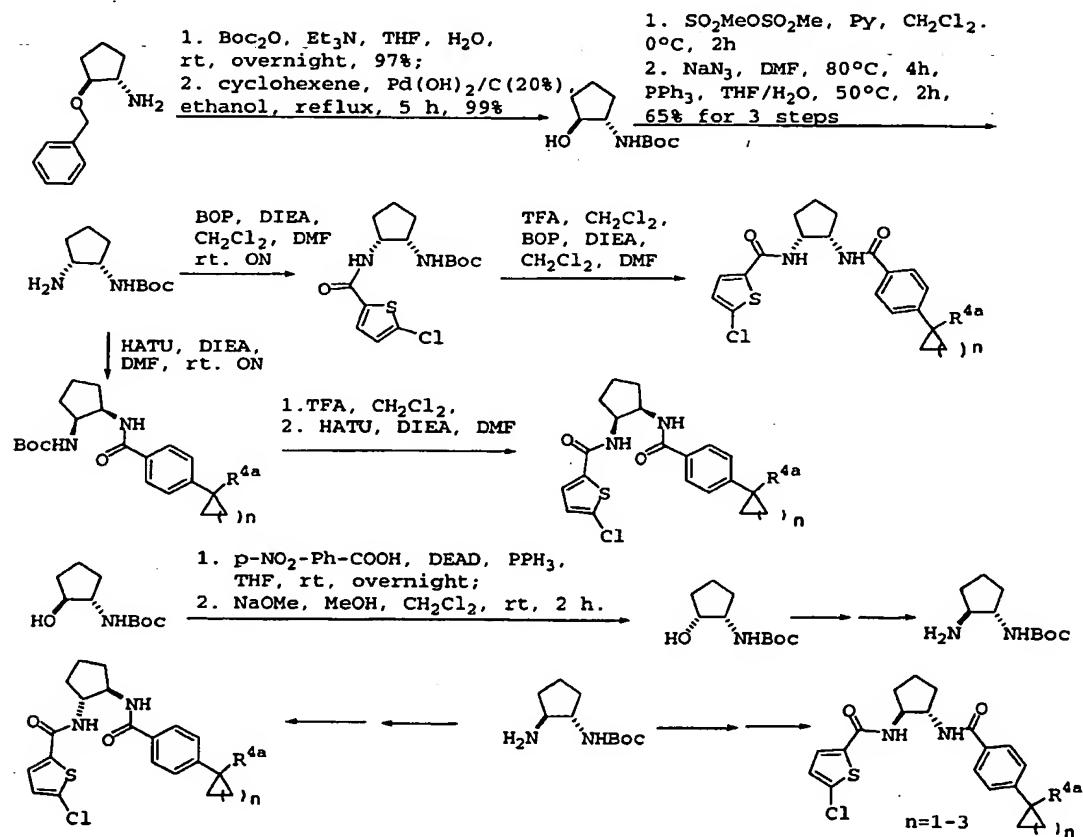
Scheme 40



Different diastereomers of compounds of the present invention can be prepared as exemplified in Scheme 41 with cyclopentyl as the central ring. Starting from

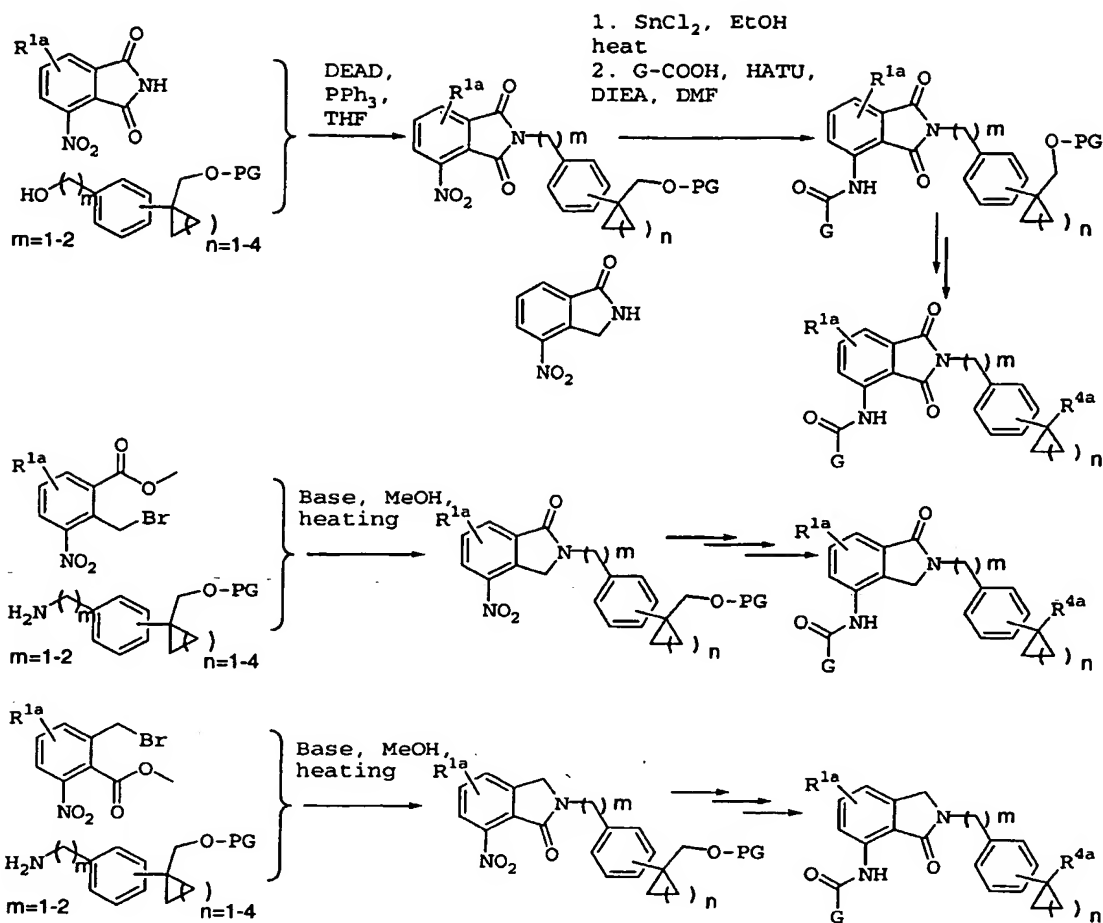
enantiomerically-pure commercially-available (1S, 2S)-2-benzyloxy-cyclopentyl amine, Boc protection followed by debenzylation gave the alcohol. SN₂ displacement with NaN₃ of the mesylate, followed by reduction of the azide afforded the key mono-boc protected diamine intermediate. Amide formation as described previously provided one pair of enantiomers. On the other hand, inversion of the stereo center of the alcohol (p-NO₂-Ph-COOH, DEAD, PPh₃, THF; then NaOMe, MeOH) followed by the same amide formation sequence should afford the other pair of enantiomers.

Scheme 41



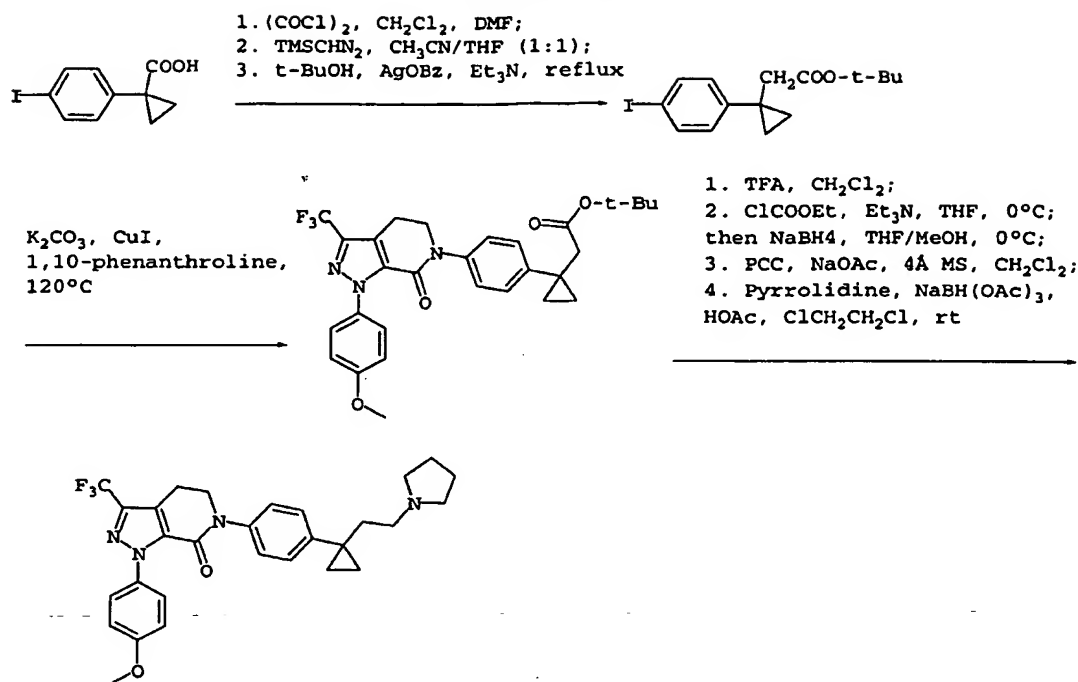
Compounds of present invention wherein M-P is an isoindole derivative can be prepared as exemplified in Scheme 42. These compounds can be obtained via standard organic transformations such as Mitsunomo reactions.

Scheme 42

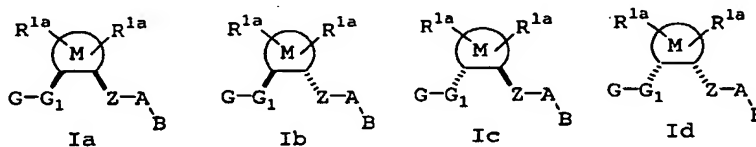


Scheme 43 depict the synthesis of compounds of the present invention wherein R^{4a} is an ethylamine derivative. This synthesis involves homologation of carboxylic acids as the key step.

Scheme 43



One diastereomer of a compound of Formula I may be more potent against fXa than the others. Thus, the following stereochemistries are considered to be a part of the present invention.



When required, separation of the racemic material can be achieved by HPLC using a chiral column or by a resolution using a resolving agent such as camphonic chloride (Steven D. Young, et al, *Antimicrobial Agents and Chemotherapy*, **1995**, 2602-2605). A chiral compound of Formula I may also be directly synthesized using a chiral catalyst or a chiral ligand (for example, Andrew S. Thompson, et al, *Tetrahedron Lett.* **1995**, 36, 8937-8940).

Other features of the invention will become apparent in the course of the following descriptions of exemplary

embodiments that are given for illustration of the invention and are not intended to be limiting thereof.

UTILITY

5 The compounds of this invention are inhibitors of factor Xa and are useful as anticoagulants for the treatment or prevention of thromboembolic disorders in mammals (i.e., factor Xa-associated disorders). In general, a thromboembolic disorder is a circulatory disease
10 caused by blood clots (i.e., diseases involving fibrin formation, platelet activation, and/or platelet aggregation). The term "thromboembolic disorders" as used herein includes arterial cardiovascular thromboembolic disorders, venous cardiovascular thromboembolic disorders,
15 and thromboembolic disorders in the chambers of the heart. The term "thromboembolic disorders" as used herein also includes specific disorders selected from, but not limited to, unstable angina or other acute coronary syndromes, first or recurrent myocardial infarction, ischemic sudden
20 death, transient ischemic attack, stroke, atherosclerosis, peripheral occlusive arterial disease, venous thrombosis, deep vein thrombosis, thrombophlebitis, arterial embolism, coronary arterial thrombosis, cerebral arterial thrombosis, cerebral embolism, kidney embolism, pulmonary embolism, and
25 thrombosis resulting from (a) prosthetic valves or other implants, (b) indwelling catheters, (c) stents, (d) cardiopulmonary bypass, (e) hemodialysis, or (f) other procedures in which blood is exposed to an artificial surface that promotes thrombosis. It is noted that
30 thrombosis includes occlusion (e.g. after a bypass) and reocclusion (e.g., during or after percutaneous transluminal coronary angioplasty). The thromboembolic disorders may result from conditions including but not
35 limited to atherosclerosis, surgery or surgical complications, prolonged immobilization, arterial fibrillation, congenital thrombophilia, cancer, diabetes,

effects of medications or hormones, and complications of pregnancy. The anticoagulant effect of compounds of the present invention is believed to be due to inhibition of factor Xa or thrombin.

5 The effectiveness of compounds of the present invention as inhibitors of factor Xa was determined using purified human factor Xa and synthetic substrate. The rate of factor Xa hydrolysis of chromogenic substrate S2222 (Diapharma/Chromogenix, West Chester, OH) was measured both
10 in the absence and presence of compounds of the present invention. Hydrolysis of the substrate resulted in the release of pNA, which was monitored spectrophotometrically by measuring the increase in absorbance at 405 nM. A decrease in the rate of absorbance change at 405 nm in the
15 presence of inhibitor is indicative of enzyme inhibition. The results of this assay are expressed as inhibitory constant, K_i .

Factor Xa determinations were made in 0.10 M sodium phosphate buffer, pH 7.5, containing 0.20 M NaCl, and 0.5% PEG 8000. The Michaelis constant, K_m , for substrate hydrolysis was determined at 25°C using the method of Lineweaver and Burk. Values of K_i were determined by allowing 0.2-0.5 nM human factor Xa (Enzyme Research Laboratories, South Bend, IN) to react with the substrate (0.20 mM-1 mM) in the presence of inhibitor. Reactions were allowed to go for 30 minutes and the velocities (rate of absorbance change vs time) were measured in the time frame of 25-30 minutes. The following relationship was used to calculate K_i values:

$$30 \quad (v_O - v_S) / v_S = I / (K_i (1 + S/K_m))$$

where:

v_0 is the velocity of the control in the absence of inhibitor:

v_s is the velocity in the presence of inhibitor;

35 I is the concentration of inhibitor;

K_i is the dissociation constant of the
enzyme:inhibitor complex;
 S is the concentration of substrate;
 K_m is the Michaelis constant.

5 Compounds tested in the above assay are considered to
be active if they exhibit a K_i of ≤ 10 μM . Preferred
compounds of the present invention have K_i 's of ≤ 1 μM .
More preferred compounds of the present invention have K_i 's
of ≤ 0.1 μM . Even more preferred compounds of the present
10 invention have K_i 's of ≤ 0.01 μM . Still more preferred
compounds of the present invention have K_i 's of ≤ 0.001 μM .
Using the methodology described above, a number of
compounds of the present invention were found to exhibit
 K_i 's of ≤ 10 μM , thereby confirming the utility of the
15 compounds of the present invention as effective Xa
inhibitors.

The antithrombotic effect of compounds of the present
invention can be demonstrated in a rabbit arterio-venous
(AV) shunt thrombosis model. In this model, rabbits
20 weighing 2-3 kg anesthetized with a mixture of xylazine (10
mg/kg i.m.) and ketamine (50 mg/kg i.m.) are used. A
saline-filled AV shunt device is connected between the
femoral arterial and the femoral venous cannulae. The AV
shunt device consists of a piece of 6-cm tygon tubing that
25 contains a piece of silk thread. Blood will flow from the
femoral artery via the AV-shunt into the femoral vein. The
exposure of flowing blood to a silk thread will induce the
formation of a significant thrombus. After forty minutes,
the shunt is disconnected and the silk thread covered with
30 thrombus is weighed. Test agents or vehicle will be given
(i.v., i.p., s.c., or orally) prior to the opening of the
AV shunt. The percentage inhibition of thrombus formation
is determined for each treatment group. The ID_{50} values
(dose which produces 50% inhibition of thrombus formation)
35 are estimated by linear regression.

The compounds of the present invention may also be useful as inhibitors of serine proteases, notably human thrombin, Factor VIIa, Factor IXa, Factor XIa, urokinase, plasma kallikrein and plasmin. Because of their inhibitory
5 action, these compounds are indicated for use in the prevention or treatment of physiological reactions, blood coagulation and inflammation, catalyzed by the aforesaid class of enzymes. Specifically, the compounds have utility as drugs for the treatment of diseases arising from
10 elevated thrombin activity such as myocardial infarction, and as reagents used as anticoagulants in the processing of blood to plasma for diagnostic and other commercial purposes.

Some compounds of the present invention were shown to
15 be direct acting inhibitors of the serine protease thrombin by their ability to inhibit the cleavage of small molecule substrates by thrombin in a purified system. *In vitro* inhibition constants were determined by the method described by Kettner et al. in *J. Biol. Chem.* **265**, 18289-
20 18297 (1990), herein incorporated by reference. In these assays, thrombin-mediated hydrolysis of the chromogenic substrate S2238 (Helena Laboratories, Beaumont, TX) was monitored spectrophotometrically. Addition of an inhibitor to the assay mixture results in decreased absorbance and is
25 indicative of thrombin inhibition. Human thrombin (Enzyme Research Laboratories, Inc., South Bend, IN) at a concentration of 0.2 nM in 0.10 M sodium phosphate buffer, pH 7.5, 0.20 M NaCl, and 0.5% PEG 6000, was incubated with various substrate concentrations ranging from 0.20 to 0.02
30 mM. After 25 to 30 minutes of incubation, thrombin activity was assayed by monitoring the rate of increase in absorbance at 405 nm that arises owing to substrate hydrolysis. Inhibition constants were derived from reciprocal plots of the reaction velocity as a function of
35 substrate concentration using the standard method of Lineweaver and Burk. Using the methodology described

above, some compounds of this invention were evaluated and found to exhibit a K_i of less than 10 μM , thereby confirming the utility of the compounds of the present invention as effective thrombin inhibitors.

5 The compounds of the present invention can be administered alone or in combination with one or more additional therapeutic agents. These include other anti-coagulant or coagulation inhibitory agents, anti-platelet or platelet inhibitory agents, thrombin inhibitors, or
10 thrombolytic or fibrinolytic agents.

 The compounds are administered to a mammal in a therapeutically effective amount. By "therapeutically effective amount" it is meant an amount of a compound of the present invention that, when administered alone or in
15 combination with an additional therapeutic agent to a mammal, is effective to prevent or ameliorate the thromboembolic disease condition or the progression of the disease.

 By "administered in combination" or "combination
20 therapy" it is meant that a compound of the present invention and one or more additional therapeutic agents are administered concurrently to the mammal being treated. When administered in combination each component may be administered at the same time or sequentially in any order
25 at different points in time. Thus, each component may be administered separately but sufficiently closely in time so as to provide the desired therapeutic effect. Other anticoagulant agents (or coagulation inhibitory agents) that may be used in combination with the compounds of this
30 invention include warfarin and heparin (either unfractionated heparin or any commercially available low molecular weight heparin), synthetic pentasaccharide, direct acting thrombin inhibitors including hirudin and argatrobanas well as other factor Xa inhibitors such as
35 those described in the publications identified above under Background of the Invention.

The term anti-platelet agents (or platelet inhibitory agents), as used herein, denotes agents that inhibit platelet function, for example by inhibiting the aggregation, adhesion or granular secretion of platelets.

5 Agents include, but are not limited to, the various known non-steroidal anti-inflammatory drugs (NSAIDs) such as aspirin, ibuprofen, naproxen, sulindac, indomethacin, mefenamate, droxicam, diclofenac, sulfinpyrazone, piroxicam, and pharmaceutically acceptable salts or

10 prodrugs thereof. Of the NSAIDs, aspirin (acetylsalicyclic acid or ASA) and piroxicam are preferred. Other suitable platelet inhibitory agents include IIb/IIIa antagonists (e.g., tirofiban, eptifibatide, and abciximab), thromboxane-A₂-receptor antagonists (e.g., ifetroban),

15 thromboxane-A₂-synthetase inhibitors, PDE-III inhibitors (e.g., dipyridamole), and pharmaceutically acceptable salts or prodrugs thereof.

The term anti-platelet agents (or platelet inhibitory agents), as used herein, is also intended to include ADP

20 (adenosine diphosphate) receptor antagonists, preferably antagonists of the purinergic receptors P₂Y₁ and P₂Y₁₂, with P₂Y₁₂ being even more preferred. Preferred P₂Y₁₂ receptor antagonists include ticlopidine and clopidogrel, including pharmaceutically acceptable salts or prodrugs thereof.

25 Clopidogrel is an even more preferred agent. Ticlopidine and clopidogrel are also preferred compounds since they are known to be gentle on the gastro-intestinal tract in use.

The term thrombin inhibitors (or anti-thrombin agents), as used herein, denotes inhibitors of the serine

30 protease thrombin. By inhibiting thrombin, various thrombin-mediated processes, such as thrombin-mediated platelet activation (that is, for example, the aggregation of platelets, and/or the granular secretion of plasminogen activator inhibitor-1 and/or serotonin) and/or fibrin

35 formation are disrupted. A number of thrombin inhibitors are known to one of skill in the art and these inhibitors

are contemplated to be used in combination with the present compounds. Such inhibitors include, but are not limited to, boroarginine derivatives, boro peptides, heparins, hirudin, argatroban, and melagatran, including pharmaceutically acceptable salts and prodrugs thereof. Boroarginine derivatives and boro peptides include N-acetyl and peptide derivatives of boronic acid, such as C-terminal α -aminoboronic acid derivatives of lysine, ornithine, arginine, homoarginine and corresponding isothiuronium analogs thereof. The term hirudin, as used herein, includes suitable derivatives or analogs of hirudin, referred to herein as hirulogs, such as disulfatohirudin.

The term thrombolytics or fibrinolytic agents (or thrombolytics or fibrinolytics), as used herein, denote agents that lyse blood clots (thrombi). Such agents include tissue plasminogen activator (natural or recombinant) and modified forms thereof, anistreplase, urokinase, streptokinase, tenecteplase (TNK), lanoteplase (nPA), factor VIIa inhibitors, PAI-1 inhibitors (i.e., inactivators of tissue plasminogen activator inhibitors), α 2-antiplasmin inhibitors, and anisoylated plasminogen streptokinase activator complex, including pharmaceutically acceptable salts or prodrugs thereof. The term anistreplase, as used herein, refers to anisoylated plasminogen streptokinase activator complex, as described, for example, in EP 028,489, the disclosure of which is hereby incorporated herein by reference herein. The term urokinase, as used herein, is intended to denote both dual and single chain urokinase, the latter also being referred to herein as prourokinase.

Examples of suitable anti-arrhythmic agents for use in combination with the present compounds include: Class I agents (such as propafenone); Class II agents (such as carvediol and propranolol); Class III agents (such as sotalol, dofetilide, amiodarone, azimilide and ibutilide); Class IV agents (such as diltiazem and verapamil); K^+

channel openers such as I_{ACh} inhibitors, and I_{Kur} inhibitors (e.g., compounds such as those disclosed in WO01/40231).

Examples of suitable anti-hypertensive agents for use in combination with the compounds of the present invention include: alpha adrenergic blockers; beta adrenergic blockers; calcium channel blockers (e.g., diltiazem, verapamil, nifedipine, amlodipine and mybefradil); diuretics (e.g., chlorothiazide, hydrochlorothiazide, flumethiazide, hydroflumethiazide, bendroflumethiazide, methylchlorothiazide, trichloromethiazide, polythiazide, benzthiazide, ethacrynic acid tricyclafen, chlorthalidone, furosemide, bumetanide, triamtrenene, amiloride, spironolactone); renin inhibitors; ACE inhibitors (e.g., captopril, zofenopril, fosinopril, enalapril, ceranopril, cilazopril, delapril, pentopril, quinapril, ramipril, lisinopril); AT-1 receptor antagonists (e.g., losartan, irbesartan, valsartan); ET receptor antagonists (e.g., sitaxsentan, atrsentan and compounds disclosed in U.S. Patent Nos. 5,612,359 and 6,043,265); Dual ET/AII antagonist (e.g., compounds disclosed in WO 00/01389); neutral endopeptidase (NEP) inhibitors; vasopepsidase inhibitors (dual NEP-ACE inhibitors) (e.g., omapatrilat, gemopatrilat and nitrates).

Examples of suitable calcium channel blockers (L-type or T-type) for use in combination with the compounds of the present invention include diltiazem, verapamil, nifedipine, amlodipine and mybefradil.

Examples of suitable cardiac glycosides for use in combination with the compounds of the present invention include digitalis and ouabain.

Examples of suitable diuretics for use in combination with the compounds of the present invention include: chlorothiazide, hydrochlorothiazide, flumethiazide, hydroflumethiazide, bendroflumethiazide, methylchlorothiazide, trichloromethiazide, polythiazide, benzthiazide, ethacrynic acid tricyclafen, chlorthalidone,

furosemide, musolimine, bumetanide, triamtrenene, amiloride, and spironolactone.

Examples of suitable mineralocorticoid receptor antagonists for use in combination with the compounds of the present invention include sprionolactone and eplirinone.

Examples of suitable phosphodiesterase inhibitors for use in combination with the compounds of the present invention include: PDE III inhibitors (such as cilostazol); and PDE V inhibitors (such as sildenafil).

Examples of suitable cholesterol/lipid lowering agents and lipid profile therapies for use in combination with the compounds of the present invention include: HMG-CoA reductase inhibitors (e.g., pravastatin, lovastatin, atorvastatin, simvastatin, fluvastatin, NK-104 (a.k.a. itavastatin, or nisvastatin or nisbastatin) and ZD-4522 (a.k.a. rosuvastatin, or atavastatin or visastatin)); squalene synthetase inhibitors; fibrates; bile acid sequestrants (such as questran); ACAT inhibitors; MTP inhibitors; lipooxygenase inhibitors; cholesterol absorption inhibitors; and cholesterol ester transfer protein inhibitors (e.g., CP-529414).

Examples of suitable anti-diabetic agents for use in combination with the compounds of the present invention include: biguanides (e.g., metformin); glucosidase inhibitors (e.g., acarbose); insulins (including insulin secretagogues or insulin sensitizers); meglitinides (e.g., repaglinide); sulfonylureas (e.g., glimepiride, glyburide and glipizide); biguanide/glyburide combinations (e.g., glucovance), thiozolidinediones (e.g., troglitazone, rosiglitazone and pioglitazone), PPAR-alpha agonists, PPAR-gamma agonists, PPAR alpha/gamma dual agonists, SGLT2 inhibitors, inhibitors of fatty acid binding protein (af2) such as those disclosed in WO00/59506, glucagon-like peptide-1 (GLP-1), and dipeptidyl peptidase IV (DP4) inhibitors.

Examples of suitable anti-depressant agents for use in combination with the compounds of the present invention include nefazodone and sertraline.

5 Examples of suitable anti-inflammatory agents for use in combination with the compounds of the present invention include: prednisone; dexamethasone; enbrel; protien tyrosine kinase (PTK) inhibitors; cyclooxygenase inhibitors (including NSAIDs, and COX-1 and/or COX-2 inhibitors); aspirin; indomethacin; ibuprofen; prioxicam; naproxen;
10 celecoxib; and/or rofecoxib.

Examples of suitable anti-osteoporosis agents for use in combination with the compounds of the present invention include alendronate and raloxifene.

15 Examples of suitable hormone replacement therapies for use in combination with the compounds of the present invention include estrogen (e.g., conjugated estrogens) and estradiol.

Examples of suitable anti-coagulants for use in combination with the compounds of the present invention
20 include heparins (e.g., unfractionated and low molecular weight heparins such as enoxaparin and dalteparin).

Examples of suitable anti-obesity agents for use in combination with the compounds of the present invention include orlistat and aP2 inhibitors (such as those
25 disclosed in WO00/59506).

Examples of suitable anti-anxiety agents for use in combination with the compounds of the present invention include diazepam, lorazepam, buspirone, and hydroxyzine pamoate.

30 Examples of suitable anti-proliferative agents for use in combination with the compounds of the present invention include cyclosporin A, paclitaxel, adriamycin; epithilones, cisplatin, and carboplatin.

35 Examples of suitable anti-ulcer and gastroesophageal reflux disease agents for use in combination with the

compounds of the present invention include famotidine, ranitidine, and omeprazole.

Administration of the compounds of the present invention (i.e., a first therapeutic agent) in combination with at least one additional therapeutic agent (i.e., a second therapeutic agent), preferably affords an efficacy advantage over the compounds and agents alone, preferably while permitting the use of lower doses of each (i.e., a synergistic combination). A lower dosage minimizes the potential of side effects, thereby providing an increased margin of safety. It is preferred that at least one of the therapeutic agents is administered in a sub-therapeutic dose. It is even more preferred that all of the therapeutic agents be administered in sub-therapeutic doses. Sub-therapeutic is intended to mean an amount of a therapeutic agent that by itself does not give the desired therapeutic effect for the condition or disease being treated. Synergistic combination is intended to mean that the observed effect of the combination is greater than the sum of the individual agents administered alone.

The compounds of the present invention are also useful as standard or reference compounds, for example as a quality standard or control, in tests or assays involving the inhibition of factor Xa. Such compounds may be provided in a commercial kit, for example, for use in pharmaceutical research involving factor Xa. For example, a compound of the present invention could be used as a reference in an assay to compare its known activity to a compound with an unknown activity. This would ensure the experimenter that the assay was being performed properly and provide a basis for comparison, especially if the test compound was a derivative of the reference compound. When developing new assays or protocols, compounds according to the present invention could be used to test their effectiveness.

The compounds of the present invention may also be used in diagnostic assays involving factor Xa. For example, the presence of factor Xa in an unknown sample could be determined by addition of chromogenic substrate S2222 to a series of solutions containing test sample and optionally one of the compounds of the present invention. If production of pNA is observed in the solutions containing test sample, but not in the presence of a compound of the present invention, then one would conclude factor Xa was present.

The present invention also encompasses an article of manufacture. As used herein, article of manufacture is intended to include, but not be limited to, kits and packages. The article of manufacture of the present invention, comprises: (a) a first container; (b) a pharmaceutical composition located within the first container, wherein the composition, comprises: a first therapeutic agent, comprising: a compound of the present invention or a pharmaceutically acceptable salt form thereof; and, (c) a package insert stating that the pharmaceutical composition can be used for the treatment of a thromboembolic disorder (as defined previously). In another embodiment, the package insert states that the pharmaceutical composition can be used in combination (as defined previously) with a second therapeutic agent to treat a thromboembolic disorder. The article of manufacture can further comprise: (d) a second container, wherein components (a) and (b) are located within the second container and component (c) is located within or outside of the second container. Located within the first and second containers means that the respective container holds the item within its boundaries.

The first container is a receptacle used to hold a pharmaceutical composition. This container can be for manufacturing, storing, shipping, and/or individual/bulk selling. First container is intended to cover a bottle,

jar, vial, flask, syringe, tube (e.g., for a cream preparation), or any other container used to manufacture, hold, store, or distribute a pharmaceutical product.

The second container is one used to hold the first container and, optionally, the package insert. Examples of the second container include, but are not limited to, boxes (e.g., cardboard or plastic), crates, cartons, bags (e.g., paper or plastic bags), pouches, and sacks. The package insert can be physically attached to the outside of the first container via tape, glue, staple, or another method of attachment, or it can rest inside the second container without any physical means of attachment to the first container. Alternatively, the package insert is located on the outside of the second container. When located on the outside of the second container, it is preferable that the package insert is physically attached via tape, glue, staple, or another method of attachment. Alternatively, it can be adjacent to or touching the outside of the second container without being physically attached.

The package insert is a label, tag, marker, etc. that recites information relating to the pharmaceutical composition located within the first container. The information recited will usually be determined by the regulatory agency governing the area in which the article of manufacture is to be sold (e.g., the United States Food and Drug Administration). Preferably, the package insert specifically recites the indications for which the pharmaceutical composition has been approved. The package insert may be made of any material on which a person can read information contained therein or thereon. Preferably, the package insert is a printable material (e.g., paper, plastic, cardboard, foil, adhesive-backed paper or plastic, etc.) on which the desired information has been formed (e.g., printed or applied).

35

Dosage and Formulation

The compounds of this invention can be administered in such oral dosage forms as tablets, capsules (each of which includes sustained release or timed release formulations), pills, powders, granules, elixirs, tinctures, suspensions, syrups, and emulsions. They may also be administered in intravenous (bolus or infusion), intraperitoneal, subcutaneous, or intramuscular form, all using dosage forms well known to those of ordinary skill in the pharmaceutical arts. They can be administered alone, but generally will be administered with a pharmaceutical carrier selected on the basis of the chosen route of administration and standard pharmaceutical practice.

The dosage regimen for the compounds of the present invention will, of course, vary depending upon known factors, such as the pharmacodynamic characteristics of the particular agent and its mode and route of administration; the species, age, sex, health, medical condition, and weight of the recipient; the nature and extent of the symptoms; the kind of concurrent treatment; the frequency of treatment; the route of administration, the renal and hepatic function of the patient, and the effect desired. A physician or veterinarian can determine and prescribe the effective amount of the drug required to prevent, counter, or arrest the progress of the thromboembolic disorder.

By way of general guidance, the daily oral dosage of each active ingredient, when used for the indicated effects, will range between about 0.001 to 1000 mg/kg of body weight, preferably between about 0.01 to 100 mg/kg of body weight per day, and most preferably between about 1.0 to 20 mg/kg/day. Intravenously, the most preferred doses will range from about 1 to about 10 mg/kg/minute during a constant rate infusion. Compounds of this invention may be administered in a single daily dose, or the total daily dosage may be administered in divided doses of two, three, or four times daily.

Compounds of this invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal routes, using transdermal skin patches. When administered in the form of a transdermal
5 delivery system, the dosage administration will, of course, be continuous rather than intermittent throughout the dosage regimen.

The compounds are typically administered in admixture with suitable pharmaceutical diluents, excipients, or
10 carriers (collectively referred to herein as pharmaceutical carriers) suitably selected with respect to the intended form of administration, that is, oral tablets, capsules, elixirs, syrups and the like, and consistent with conventional pharmaceutical practices.

15 For instance, for oral administration in the form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic, pharmaceutically acceptable, inert carrier such as lactose, starch, sucrose, glucose, methyl cellulose, magnesium stearate, dicalcium
20 phosphate, calcium sulfate, mannitol, sorbitol and the like; for oral administration in liquid form, the oral drug components can be combined with any oral, non-toxic, pharmaceutically acceptable inert carrier such as ethanol, glycerol, water, and the like. Moreover, when desired or
25 necessary, suitable binders, lubricants, disintegrating agents, and coloring agents can also be incorporated into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia,
30 tragacanth, or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes, and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride, and the like. Disintegrators
35 include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum, and the like.

The compounds of the present invention can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines.

Compounds of the present invention may also be coupled with soluble polymers as targetable drug carriers. Such polymers can include polyvinylpyrrolidone, pyran copolymer, polyhydroxypropylmethacrylamide-phenol, polyhydroxyethylaspartamidephenol, or polyethyleneoxide-polylysine substituted with palmitoyl residues. Furthermore, the compounds of the present invention may be coupled to a class of biodegradable polymers useful in achieving controlled release of a drug, for example, polylactic acid, polyglycolic acid, copolymers of polylactic and polyglycolic acid, polyepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihdropyrans, polycyanoacylates, and crosslinked or amphipathic block copolymers of hydrogels.

Dosage forms (pharmaceutical compositions) suitable for administration may contain from about 1 milligram to about 100 milligrams of active ingredient per dosage unit. In these pharmaceutical compositions the active ingredient will ordinarily be present in an amount of about 0.5-95% by weight based on the total weight of the composition.

Gelatin capsules may contain the active ingredient and powdered carriers, such as lactose, starch, cellulose derivatives, magnesium stearate, stearic acid, and the like. Similar diluents can be used to make compressed tablets. Both tablets and capsules can be manufactured as sustained release products to provide for continuous release of medication over a period of hours. Compressed tablets can be sugar coated or film coated to mask any unpleasant taste and protect the tablet from the

atmosphere, or enteric coated for selective disintegration in the gastrointestinal tract.

Liquid dosage forms for oral administration can contain coloring and flavoring to increase patient
5 acceptance.

In general, water, a suitable oil, saline, aqueous dextrose (glucose), and related sugar solutions and glycols such as propylene glycol or polyethylene glycols are suitable carriers for parenteral solutions. Solutions for
10 parenteral administration preferably contain a water soluble salt of the active ingredient, suitable stabilizing agents, and if necessary, buffer substances. Antioxidizing agents such as sodium bisulfite, sodium sulfite, or ascorbic acid, either alone or combined, are suitable
15 stabilizing agents. Also used are citric acid and its salts and sodium EDTA. In addition, parenteral solutions can contain preservatives, such as benzalkonium chloride, methyl-or propyl-paraben, and chlorobutanol.

Suitable pharmaceutical carriers are described in
20 Remington's Pharmaceutical Sciences, Mack Publishing Company, a standard reference text in this field.

Representative useful pharmaceutical dosage-forms for administration of the compounds of this invention can be illustrated as follows:

25 Capsules

A large number of unit capsules can be prepared by filling standard two-piece hard gelatin capsules each with 100 milligrams of powdered active ingredient, 150 milligrams of lactose, 50 milligrams of cellulose, and 6
30 milligrams magnesium stearate.

Soft Gelatin Capsules

A mixture of active ingredient in a digestible oil such as soybean oil, cottonseed oil or olive oil may be prepared and injected by means of a positive displacement
35 pump into gelatin to form soft gelatin capsules containing

100 milligrams of the active ingredient. The capsules should be washed and dried.

Tablets

Tablets may be prepared by conventional procedures so that the dosage unit is 100 milligrams of active ingredient, 0.2 milligrams of colloidal silicon dioxide, 5 milligrams of magnesium stearate, 275 milligrams of microcrystalline cellulose, 11 milligrams of starch and 98.8 milligrams of lactose. Appropriate coatings may be applied to increase palatability or delay absorption.

Injectable

A parenteral composition suitable for administration by injection may be prepared by stirring 1.5% by weight of active ingredient in 10% by volume propylene glycol and water. The solution should be made isotonic with sodium chloride and sterilized.

Suspension

An aqueous suspension can be prepared for oral administration so that each 5 mL contain 100 mg of finely divided active ingredient, 200 mg of sodium carboxymethyl cellulose, 5 mg of sodium benzoate, 1.0 g of sorbitol solution, U.S.P, and 0.025 mL of vanillin.

Where the compounds of this invention are combined with other anticoagulant agents, for example, a daily dosage may be about 0.1 to 100 milligrams of the compound of Formula I and about 1 to 7.5 milligrams of the second anticoagulant, per kilogram of patient body weight. For a tablet dosage form, the compounds of this invention generally may be present in an amount of about 5 to 10 milligrams per dosage unit, and the second anti-coagulant in an amount of about 1 to 5 milligrams per dosage unit.

Where the compounds of the present invention are administered in combination with an anti-platelet agent, by way of general guidance, typically a daily dosage may be about 0.01 to 25 milligrams of the compound of Formula I and about 50 to 150 milligrams of the anti-platelet agent,

preferably about 0.1 to 1 milligrams of the compound of Formula I and about 1 to 3 milligrams of antiplatelet agents, per kilogram of patient body weight.

Where the compounds of Formula I are administered in combination with thrombolytic agent, typically a daily dosage may be about 0.1 to 1 milligrams of the compound of Formula I, per kilogram of patient body weight and, in the case of the thrombolytic agents, the usual dosage of the thrombolytic agent when administered alone may be reduced by about 70-80% when administered with a compound of Formula I.

Where two or more of the foregoing second therapeutic agents are administered with the compound of Formula I, generally the amount of each component in a typical daily dosage and typical dosage form may be reduced relative to the usual dosage of the agent when administered alone, in view of the additive or synergistic effect of the therapeutic agents when administered in combination.

Particularly when provided as a single dosage unit, the potential exists for a chemical interaction between the combined active ingredients. For this reason, when the compound of Formula I and a second therapeutic agent are combined in a single dosage unit they are formulated such that although the active ingredients are combined in a single dosage unit, the physical contact between the active ingredients is minimized (that is, reduced). For example, one active ingredient may be enteric coated. By enteric coating one of the active ingredients, it is possible not only to minimize the contact between the combined active ingredients, but also, it is possible to control the release of one of these components in the gastrointestinal tract such that one of these components is not released in the stomach but rather is released in the intestines. One of the active ingredients may also be coated with a material that affects a sustained-release throughout the gastrointestinal tract and also serves to minimize physical

contact between the combined active ingredients.
Furthermore, the sustained-released component can be additionally enteric coated such that the release of this component occurs only in the intestine. Still another
5 approach would involve the formulation of a combination product in which the one component is coated with a sustained and/or enteric release polymer, and the other component is also coated with a polymer such as a low-viscosity grade of hydroxypropyl methylcellulose (HPMC) or
10 other appropriate materials as known in the art, in order to further separate the active components. The polymer coating serves to form an additional barrier to interaction with the other component.

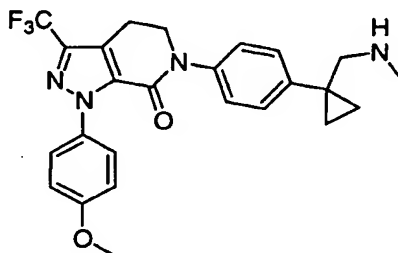
These as well as other ways of minimizing contact
15 between the components of combination products of the present invention, whether administered in a single dosage form or administered in separate forms but at the same time by the same manner, will be readily apparent to those skilled in the art, once armed with the present disclosure.

20 Other features of the invention will become apparent in the course of the following descriptions of exemplary embodiments that are afforded for illustration of the invention and are not intended to be limiting thereof.

25

Example 1

**1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopropyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one, trifluoroacetic acid salt**



30

Part A. δ -Valerolactam (22.22 g, 222.89 mmol) was stirred in CHCl_3 (500 mL) at 0°C . PCl_5 (140.0 g, 68.29 mmol) was added portionwise. The resulting slurry was stirred at reflux for 3 h until the solution became clear. The mixture was cooled in an ice bath and H_2O was added carefully until the PCl_5 was quenched completely. The two layers were separated. The organic layer was washed with H_2O (3x) and brine (2x), dried over MgSO_4 , filtered, and concentrated to dryness to give 3,3-dichloro-2-piperidinone (31.63 g, yield: 85%). This solid (16.50 g, 98.80 mmol) was dissolved in DMF (20 mL), and Li_2CO_3 (21.93 g, 296.40 mmol, 3.0 eq) was added. The mixture was stirred at 120°C for 1 day. The solvent was further concentrated, and 1N HCl was added to acidify the mixture. It was then extracted with CHCl_3 (6x). The organic layers were washed with H_2O , brine, dried over MgSO_4 , and concentrated to dryness to give almost pure 3-chloro-5,6-dihydro-2(1H)-pyridinone (11.13 g, 87%).

Part B. The product from Part A (5.50 g, 41.98 mmol) and 2,2,2-trifluoro-N-(4-methoxyphenyl)-ethanehydrazonoyl bromide (12.90 g, 43.58 mmol) were stirred in toluene (100 mL) at room temperature under N_2 . Et_3N (28.0 mL, 200.1 mmol) was then added. The mixture was stirred at 85°C for 15 h. It was cooled to room temperature and extracted with EtOAc (3x). The organic layers were washed with H_2O (2x) and brine (2x), dried over MgSO_4 , filtered, and concentrated to dryness. The residue was purified by flash column chromatography (silica gel, CH_2Cl_2 , then $\text{CH}_2\text{Cl}_2:\text{EtOAc}=4:1$, then EtOAc) to produce 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one as a light-tan solid (5.09 g, yield: 39%). LC/MS (ESI⁺) 312.4 (M+H)⁺.

Part C. 1-Phenyl-cyclopropylcarboxylic acid (16.50 g, 101.0 mmol) was stirred in HOAc (70 mL) at RT under N₂. I₂ (27.94 g, 101.0 mmol) was added, followed by the addition of NaIO₃ (4.98 g, 25.25 mmol) and conc. H₂SO₄ (1 mL). The
5 resulting mixture was stirred at 70°C for 3 days. LC-MS showed completion of the reaction. The cooled mixture was concentrated, poured into H₂O, and extracted with EtOAc (2x). The organic layer was washed with sodium thiosulfate (2x) and brine, dried over MgSO₄, filtered, and
10 concentrated to dryness to give almost pure 4-iodophenylcyclopropyl carboxylic acid (23.56 g, yield: 81%). LC/MS (ESI⁺) 472.4 (M+H)⁺.

Part D. The product of part C (0.22 g, 0.76 mmol) and the
15 product from part B (0.11 g, 0.35 mmol) were stirred in DMSO (0.5 mL) under N₂. K₂CO₃ (0.15 g, 1.09 mmol, 3.0 eq) was added, followed by the addition of 1,10-phenanthroline (28 mg, 20mol%) and CuI (30 mg, 20%mol). The resulting mixture was stirred at 130°C overnight. LC-MS showed
20 completion of the reaction. EtOAc was added to the cooled solution. It was washed with 1N HCl, H₂O, and brine; dried over MgSO₄; filtered; and concentrated in vacuo to give almost pure 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-
25 c]pyridin-6-yl] phenyl}cyclopropanecarboxylic acid (460 mg, 94%). LC/MS (ESI⁺) 472.6 (M+H)⁺, t_R = 2.39 min (35-98% CH₃CN in H₂O in a 6-min run).

Part E. The product from Part D (0.28 g, 0.59 mmol) was
30 stirred in THF (5 mL) at 0°C under N₂. Et₃N (0.15 mL, 1.06 mmol) was added, followed by dropwise addition of ClCO₂Et (0.098 mL, 1.03 mmol). The reaction mixture was then stirred at 0°C for 1 h. TLC showed the completion of the reaction. The mixture was filtered through a filter funnel

and rinsed with anhydrous THF. The THF filtrate (ca. 10 mL) was stirred at 0°C under N₂. MeOH (2.5 mL) was added followed by addition of NaBH₄ (0.31 g, 8.16 mmol, 13.8 eq) portionwise. The resulting mixture was stirred at 0°C for 5 15 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give 6-{4-[1-
10 (hydroxymethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.23 g, 84.7%). LC/MS(ESI⁺) 458.6 (M+H)⁺, t_R = 6.06 min (5-98% CH₃CN in H₂O in a 10-min run). ¹H NMR (CHCl₃) δ 7.38 (d, J = 9.1 Hz, 2H), 7.29 (d, J = 8.4 Hz,
15 2H), 7.16 (m, 2H), 6.83 (d, J = 9.1 Hz, 2H), 4.03 (t, J = 6.6 Hz, 2H), 3.72 (s, 3H), 3.54 (s, 2H), 3.07 (t, J = 6.6 Hz, 2H), 0.75 (s, br, 2H) ppm.

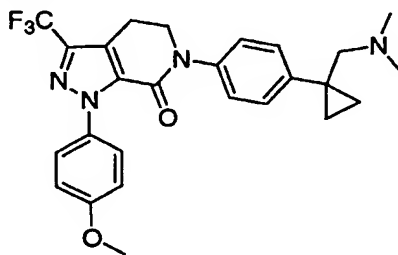
The above aldehyde (0.22 g, 0.49 mmol) was stirred in
20 anhydrous CH₂Cl₂ (5 mL) at RT under N₂. NaOAc (88 mg, 1.07 mmol, 2.2 eq) and 4Å molecular sieves (200 mg) were added, followed by the addition of PCC (0.19 g, 0.88 mmol, 1.8 eq). The resulting slurry was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The
25 mixture was filtered through Celite® and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give almost pure 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-
30 c]pyridin-6-yl]phenyl}cyclopropanecarbaldehyde (0.20 g, yield: 87.4%). LC/MS (ESI⁺) 455.4 (M+H)⁺.

Part F. The product from Part E (20 mg, 0.95 mmol), methylamine hydrochloride (20 mg, excess) were stirred in

dichloroethane (0.7 mL) in a capped vial. $\text{NaBH}(\text{OAc})_3$ (50 mg, excess) was added, followed by addition of one drop of HOAc. The reaction mixture was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was evaporated, and dissolved in aqueous MeOH. It was then purified by prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run, $t_R = 4.65$ min) to obtain the product 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (18 mg, 86%). LC/MS (ESI⁺) 471.4 (M+H)⁺. HRMS $\text{C}_{25}\text{H}_{26}\text{O}_2\text{F}_3\text{N}_4$ (M+H)⁺ 471.2011 calcd for 471.2008. ¹H NMR (acetone-*d*₆) δ 7.51 (m, 4H), 7.32 (d, $J = 8.5$ Hz, 2H), 6.99 (d, $J = 9.1$ Hz, 2H), 4.16 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.37 (m, 2H), 3.16 (t, $J = 6.3$ Hz, 2H), 2.70 (s, 3H), 1.27 (d, $J = 6.2$ Hz, 6H), 1.12 (m, 2H), 0.96 (m, 2H) ppm.

Example 2

6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

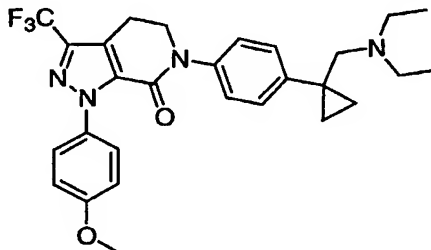


Following a procedure analogous to that used for step F in Example 1, but using dimethylamine hydrochloride, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run). LC/MS (ESI⁺) 485.4 (M+H)⁺, $t_R = 4.66$ min. HRMS $\text{C}_{26}\text{H}_{28}\text{O}_2\text{F}_3\text{N}_4$ (M+H)⁺ 485.2158 calcd for 485.2164. ¹H NMR (acetone-*d*₆) δ 7.51 (m, 4H), 7.35 (d, $J = 8.4$ Hz, 2H), 6.98 (d, $J = 9.0$ Hz, 2H), 4.17 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.60 (m, 2H), 3.16 (t, J

= 6.3 Hz, 2H), 2.82 (s, 6H), 1.15 (m, 2H), 1.08 (m, 2H) ppm.

Example 3

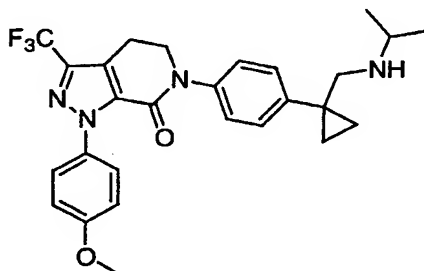
5 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



Following a procedure analogous to that used for step F in
10 Example 1, but using diethylamine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 513.4 (M+H)⁺, t_R = 4.79 min.

15 Example 4

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



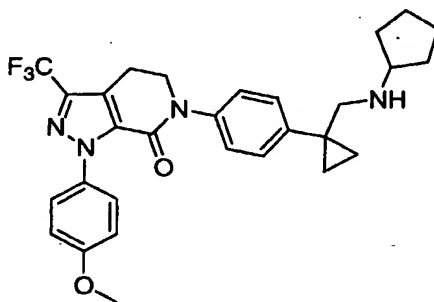
20 Following a procedure analogous to that used for step F in Example 1, but using isopropylamine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 499.4 (M+H)⁺, t_R = 4.79 min. ¹H NMR (acetone-d₆) δ 7.51 (d, J = 8.4 Hz, 4H),
25 7.31 (d, J = 8.1 Hz, 2H), 6.99 (d, J = 8.4 Hz, 2H), 4.16

(t, J = 6.5 Hz, 2H), 3.83 (s, 3H), 3.45 (m, 1H), 3.37 (m, 2H), 3.16 (t, J = 6.3 Hz, 2H), 1.27 (d, J = 6.2 Hz, 6H), 1.13 (m, 2H), 0.93 (m, 2H) ppm.

5

Example 5

6-(4-{1-[(cyclopentylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



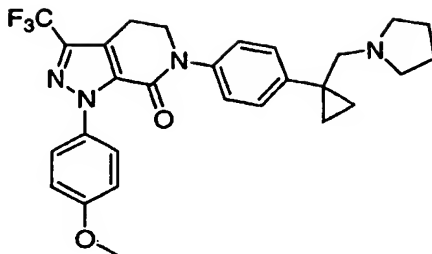
10 Following a procedure analogous to that used for step F in Example 1, but using cyclopentylamine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 525.4 (M+H)⁺, t_R = 5.03 min. HRMS C₂₉H₃₂O₂F₃N₄ (M+H)⁺ 525.2486
 15 calcd for 525.2477. ¹H NMR (acetone-*d*₆) δ 7.50 (m, 4H), 7.31 (d, J = 8.0 Hz, 2H), 6.99 (d, J = 8.8 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.62 (m, 1H), 3.31 (m, 2H), 3.16 (t, J = 6.5 Hz, 2H), 1.99 (m, 2H), 1.68 (m, 4H), 1.50 (m, 2H), 1.14 (m, 2H), 0.96 (m, 2H) ppm.

20

Example 6

1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

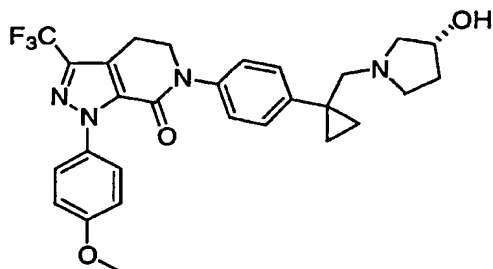
25



Following a procedure analogous to that used for step F in Example 1, but using pyrrolidine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98%
 5 CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 511.4 (M+H)⁺, t_R = 4.86 min. HRMS C₂₈H₃₀O₂F₃N₄ (M+H)⁺ 511.2320 calcd for 511.2321. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.36 (d, *J* = 8.4 Hz, 2H), 6.98 (d, *J* = 8.8 Hz, 2H), 4.17 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.60 (m, 2H), 3.49 (m, 2H), 3.16 (t, *J* =
 10 6.3 Hz, 2H), 2.93 (m, 2H), 1.93 (m, 4H), 1.14 (m, 2H), 1.01 (m, 2H) ppm.

Example 7

6-[4-(1-((3*R*)-3-hydroxy-1-pyrrolidinyl)methyl)cyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one, trifluoroacetic acid salt
 15



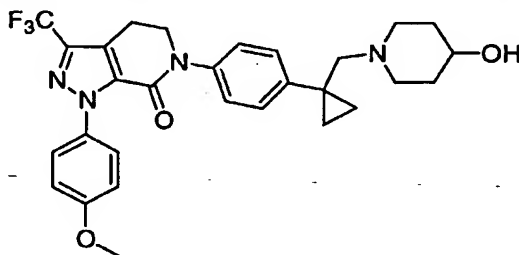
Following a procedure analogous to that used for step F in
 20 Example 1, but using (*R*)-(+)-pyrrolidinol, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 527.4 (M+H)⁺, t_R = 4.49 min. ¹H NMR (acetone-*d*₆) δ 7.55 (d, *J* = 8.4 Hz, 2H), 7.50 (d, *J* = 9.2 Hz, 2H), 7.36 (d, *J* = 8.1 Hz,

2H), 6.98 (d, $J = 9.1$ Hz, 2H), 4.17 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.72 (m, 1H), 3.40-3.27 (m, 4H), 3.16 (t, $J = 6.3$ Hz, 2H), 1.92 (m, 2H), 1.15 (m, 2H), 1.02 (m, 2H) ppm.

5

Example 8

6-(4-{1-[(4-hydroxy-1-piperidinyl)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



10

Following a procedure analogous to that used for step F in Example 1, but using 4-hydroxypiperidine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 541.4

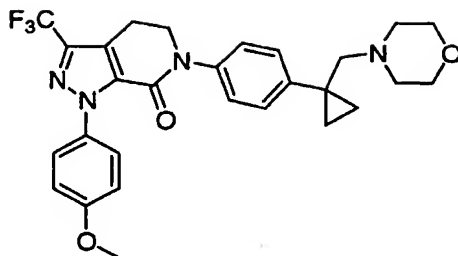
15 (M+H)⁺, $t_R = 4.63$ min. ¹H NMR (acetone-*d*₆) δ 7.51 (m, 4H), 7.33 (d, $J = 8.4$ Hz, 2H), 6.98 (d, $J = 9.1$ Hz, 2H), 4.17 (t, $J = 6.5$ Hz, 2H), 3.83 (s, 3H), 3.58 (m, 2H), 3.38 (m, 5H), 3.16 (t, $J = 6.3$ Hz, 2H), 1.92 (m, 2H), 1.78 (m, 2H), 1.19 (m, 2H), 1.05 (m, 2H) ppm.

20

Example 9

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

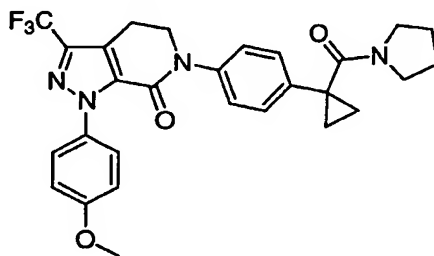
25



Following a procedure analogous to that used for step F in Example 1, but using morpholine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98%
 5 CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 527.4 (M+H)⁺, t_R = 6.08 min. HRMS C₂₈H₃₀O₃F₃N₄ (M+H)⁺ 527.2280 calcd for 527.2270. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.33 (d, *J* = 8.4 Hz, 2H), 6.98 (d, *J* = 9.1 Hz, 2H), 4.17 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.64 (m, 1H), 3.17 (t, *J* = 6.6 Hz, 2H),
 10 1.92 (m, 2H), 1.18 (t, *J* = 4.4 Hz, 2H), 1.06 (t, *J* = 4.4 Hz, 2H) ppm.

Example 10

1- (4-methoxyphenyl)-6- (4- [1- (1-
 15 pyrrolidinylcarbonyl)cyclopropyl]phenyl)-3-
 (trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
 c]pyridin-7-one

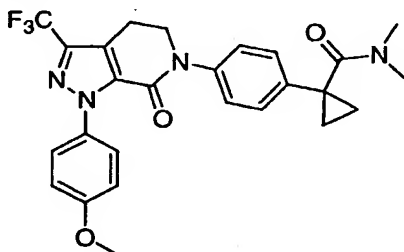


Part A. The product from part D of Example 1 (0.21 g, 0.45
 20 mmol) was stirred in CH₂Cl₂ (10 mL) at RT under N₂. SOCl₂ (0.1 mL) was added. The mixture was stirred at RT for 1 h. It was then concentrated to dryness in vacuo. The product (20 mg) was stirred in CH₂Cl₂ (0.6 mL). Pyrrolidine (0.02 mL) was added, followed by the addition of DIEA (0.05 mL)
 25 and one piece of DMAP. The mixture was stirred at RT for

0.5 h. LC-MS showed completion of the reaction. It was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to afford 1-(4-methoxyphenyl)-6-(4-[1-(1-pyrrolidinyl carbonyl)cyclopropyl]phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (15 mg, yield, 67%). LC/MS(ESI⁺) 525.4 (M+H)⁺, t_R = 6.17 min. ¹H NMR (acetone-d₆) δ 7.49 (d, J = 9.2 Hz, 2H), 7.31 (d, J = 8.8 Hz, 2H), 7.20 (d, J = 8.5 Hz, 2H), 6.97 (d, J = 9.1 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.35 (m, 2H), 3.20 (m, 2H), 3.16 (t, J = 6.6 Hz, 2H), 1.74 (m, 4H), 1.31 (m, 2H), 1.08 (m, 2H) ppm.

Example 11

1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-N,N-dimethylcyclopropanecarboxamide



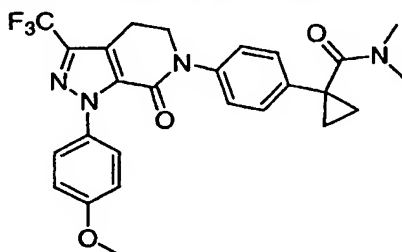
Following a procedure analogous to that used for the preparation of Example 10, but using dimethylamine hydrochloride, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 499.4 (M+H)⁺, t_R = 6.01 min. HRMS C₂₆H₂₆O₃F₃N₄ (M+H)⁺ 499.1948 calcd for 499.1957. ¹H NMR (acetone-d₆) δ 7.49 (d, J = 9.1 Hz, 2H), 7.32 (d, J = 8.8 Hz, 2H), 7.17 (d, J = 8.8 Hz, 2H), 6.97 (d, J = 8.8 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.16 (t, J = 6.2 Hz, 2H), 2.87 (s, 6H), 1.30 (m, 2H), 1.14 (m, 2H) ppm.

Example 12

0.5 h. LC-MS showed completion of the reaction. It was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to afford 1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinyl carbonyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (15 mg, yield, 67%). LC/MS(ESI⁺) 525.4 (M+H)⁺, t_R = 6.17 min. ¹H NMR (acetone-d₆) δ 7.49 (d, J = 9.2 Hz, 2H), 7.31 (d, J = 8.8 Hz, 2H), 7.20 (d, J = 8.5 Hz, 2H), 6.97 (d, J = 9.1 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.35 (m, 2H), 3.20 (m, 2H), 3.16 (t, J = 6.6 Hz, 2H), 1.74 (m, 4H), 1.31 (m, 2H), 1.08 (m, 2H) ppm.

Example 11

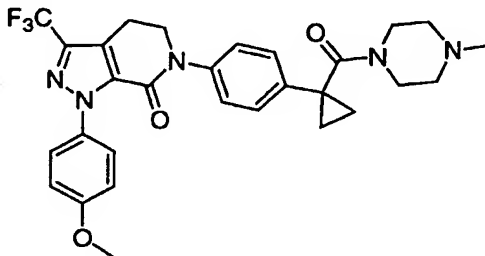
15 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-
N,N-dimethylcyclopropanecarboxamide



Following a procedure analogous to that used for the preparation of Example 10, but using dimethylamine hydrochloride, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 499.4 (M+H)⁺, t_R = 6.01 min. HRMS C₂₆H₂₆O₃F₃N₄ (M+H)⁺ 499.1948 calcd for 499.1957. ¹H NMR (acetone-d₆) δ 7.49 (d, J = 9.1 Hz, 2H), 7.32 (d, J = 8.8 Hz, 2H), 7.17 (d, J = 8.8 Hz, 2H), 6.97 (d, J = 8.8 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.16 (t, J = 6.2 Hz, 2H), 2.87 (s, 6H), 1.30 (m, 2H), 1.14 (m, 2H) ppm.

Example 12

1-(4-methoxyphenyl)-6-(4-{1-[(4-methyl-1-piperazinyl)carbonyl]cyclopropyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



5

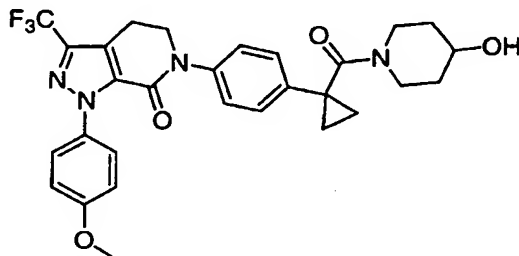
Following a procedure analogous to that used for the preparation of Example 10, but using 4-methylpiperazine, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

- 10 LC/MS(ESI⁺) 554.6 (M+H)⁺, *t_R* = 4.49 min. HRMS C₂₉H₃₁O₃F₃N₅ (M+H)⁺ 554.2384 calcd for 554.2379. ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J* = 8.8 Hz, 2H), 7.30 (AA'BB', *J* = 8.4 Hz, 4H), 6.97 (d, *J* = 9.1 Hz, 2H), 4.17 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.16 (t, *J* = 6.4 Hz, 2H), 2.85 (s, 3H), 1.39 (m, 15 2H), 1.20 (m, 2H) ppm.

Example 13

6-(4-[1-(4-hydroxypiperidine-1-carbonyl)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

20



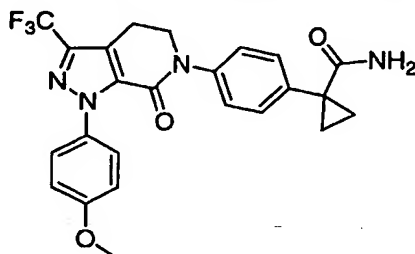
- Following a procedure analogous to that used for the preparation of Example 10, but using morpholine, the title compound was prepared. The product was purified by RP-prep
- 25

LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 555.4 (M+H)⁺, *t_R* = 5.50 min. HRMS C₂₉H₃₀O₄F₃N₄ (M+H)⁺ 555.2241 calcd for 555.2219.

5

Example 14

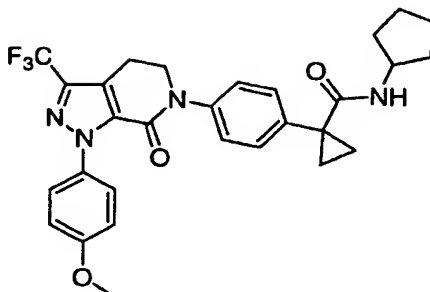
1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropanecarboxamide



- 10 Following a procedure analogous to that used for the preparation of Example 10, but using concentrated NH₄OH as the amine source and THF as solvent, the title compound was prepared. The product was purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 471.6 (M+H)⁺, *t_R* =
- 15 2.56 min (10-90% CH₃CN/H₂O in a 10-min run). ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J* = 8.8 Hz, 2H), 7.43 (d, *J* = 8.8 Hz, 2H), 7.26 (d, *J* = 8.8 Hz, 2H), 6.98 (d, *J* = 8.8 Hz, 2H), 4.18 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.17 (t, *J* = 6.6 Hz, 2H), 1.40 (m, 2H), 0.96 (m, 2H) ppm. ¹⁹F NMR
- 20 (acetone-*d*₆) δ -77.14 ppm.

Example 15

- 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropanecarboxylic acid cyclopentylamide**
- 25

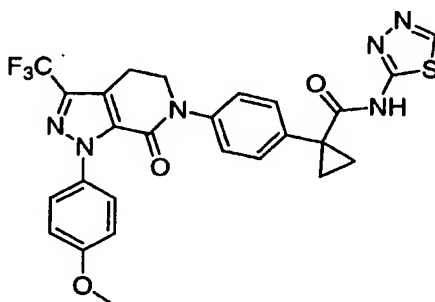


Following a procedure analogous to that used for the preparation of Example 10, but using cyclopentylamine, the title compound was prepared. The product was purified by
 5 RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS(ESI⁺) 539.4 (M+H)⁺, t_R = 6.65 min. ¹H NMR (acetone-d₆) δ 7.51 (d, J = 8.8 Hz, 2H), 7.37 (m, 4H), 6.99 (d, J = 8.8 Hz, 2H), 4.16 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 4.05 (m, 1H), 3.16 (t, J = 6.2 Hz, 2H), 1.79 (m, 2H), 1.45 (m,
 10 4H), 1.22 (m, 2H), 1.39 (m, 2H), 0.92 (m, 2H) ppm.

Example 16

1-**{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-**
 15 **N-(1,3,4-thiadiazol-2-yl)cyclopropanecarboxamide**



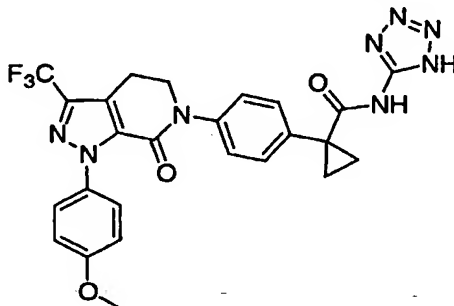
Following a procedure analogous to that used for the preparation of Example 10, but using 2-aminothiadiazole, the title compound was prepared. The product was purified by
 20 by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS(ESI⁺) 555.4 (M+H)⁺, t_R = 6.17 min. ¹H NMR (acetone-d₆) δ 8.96 (s, 1H), 7.54 (m, 4H), 7.43 (d, J = 8.4 Hz, 2H), 6.98 (d, J = 8.9 Hz, 2H), 4.22 (t, J = 6.6 Hz, 2H), 3.84

(s, 3H), 3.16 (t, J = 6.2 Hz, 2H), 1.67 (m, 2H), 1.31 (m, 2H) ppm.

Example 17

5 **1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-
N-(1H-tetrazol-5-yl)cyclopropanecarboxamide**

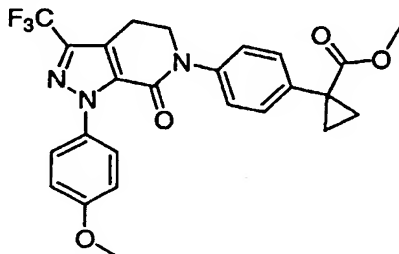


10 Following a procedure analogous to that used for the
preparation of Example 10, but using 5-amino-1H-tetrazole,
the title compound was prepared. The product was purified
by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS(ESI⁺) 539.6 (M+H), t_R = 5.86 min. ¹H NMR (acetone-*d*₆)
δ 7.53 (m, 4H), 7.39 (d, J = 8.4 Hz, 2H), 6.99 (d, J = 9.1
15 Hz, 2H), 4.20 (t, J = 6.6 Hz, 2H), 3.84 (s, 3H), 3.18 (t, J
= 6.6 Hz, 2H), 1.68 (m, 2H), 1.29 (m, 2H) ppm.

Example 18

20 **methyl 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl)cyclopropanecarboxylate**



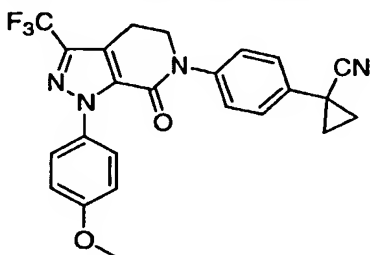
The product from part D in Example 1 (mg, mmol) was stirred
in anhydrous MeOH (5 mL) at RT. Catalytic amount of conc.

HCl was added. The resulting solution was stirred at RT overnight. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS(ESI⁺) 486.6 (M+H)⁺, t_R = 2.98 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR

(acetone-*d*₆) δ 7.50 (d, *J* = 8.8 Hz, 2H), 7.35 (AA'BB', *J* = 8.8 Hz, 4H), 6.97 (d, *J* = 9.1 Hz, 2H), 4.18 (t, *J* = 6.6 Hz, 2H), 3.82 (s, 3H), 3.55 (s, 3H), 3.17 (t, *J* = 6.4 Hz, 2H), 1.49 (m, 2H), 1.16 (m, 2H) ppm.

Example 19

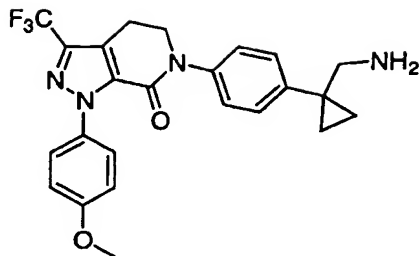
1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropanecarbonitrile



The product from Example 14 (22 mg, 0.047 mmol) was stirred in DMF (0.3 mL) at RT in a capped vial. SOCl₂ (0.05 mL) was added. The mixture was stirred at RT for 1.5 h. LC-MS showed completion of the reaction. Prep LC-MS purification (35-98% CH₃CN in H₂O) provided the title compound (15 mg, yield, 71%). LC/MS(ESI⁺) 453.4 (M+H)⁺, t_R = 5.24 min. ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J* = 9.2 Hz, 2H), 7.39 (m, 4H), 6.97 (d, *J* = 8.8 Hz, 2H), 4.18 (t, *J* = 6.6 Hz, 2H), 3.84 (s, 3H), 3.17 (t, *J* = 6.6 Hz, 2H), 1.71 (m, 2H), 1.48 (m, 2H) ppm. ¹⁹F NMR (acetone-*d*₆) δ -77.16 ppm.

Example 20

6-{4-[1-(aminomethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-*c*]pyridin-7-one, trifluoroacetic acid salt



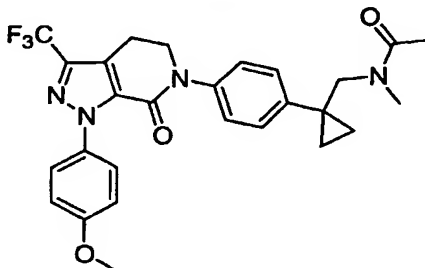
Part A. The product from part E in Example 1 (24 mg, 0.052 mmol) was stirred in CH_2Cl_2 (1 mL) at 0°C under N_2 . Et_3N (11 μL , 1.5 eq) was added followed by the dropwise addition of MsCl (4.5 μL , 1.1 eq). The mixture was stirred at 0°C for 1 h. TLC showed completion of the reaction. Sat'd NH_4Cl was added. The mixture was extracted with EtOAc . The organic layer was washed with H_2O and brine, dried over Na_2SO_4 , filtered, and concentrated to dryness. The residue was dissolved in DMF (1 mL). NaN_3 (50 mg, mmol) was added. The mixture was stirred at RT under N_2 overnight. LC-MS showed the azide as the major component in the mixture. Sat'd NH_4Cl was then added. The mixture was extracted with EtOAc . And the organic layer was washed with H_2O and brine, dried over Na_2SO_4 , filtered, and concentrated to dryness to give crude 6-(4-[1-azidomethylcyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one. LC/MS(ESI⁺) 483.4 (M+H)⁺, t_R = 3.06 min (10-90% CH_3CN in H_2O in a 4-min run).

Part B. The product from part A (18 mg) and PPh_3 (38 mg) were stirred in THF (1.5 mL) at RT for 20 min. H_2O (0.3 mL) was added, and the mixture was stirred at 30°C for 2 h. The solvents were evaporated. The residue was purified by prep LC-MS (5-98% CH_3CN in H_2O) to give pure 6-(4-[1-(aminomethyl)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-

c]pyridin-7-one (7 mg, yield: 29%). LC/MS (ESI⁺) 457.4 (M+H)⁺.

Example 21

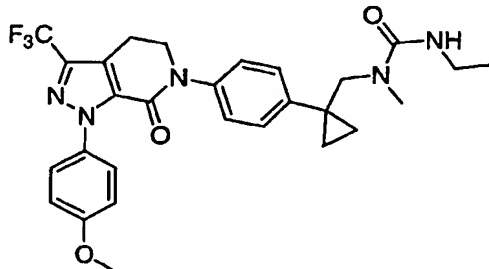
5 ***N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylacetamide**



The product of Example 1 (40 mg, 0.084 mmol) was stirred in
 10 CH₂Cl₂ (1 mL) in a capped vial at RT. Et₃N (4 drops) was added followed by addition of acetyl chloride (2 drops). The resulting mixture was stirred at RT for 10 min. LC-MS showed completion of the reaction. After evaporation of the solvents, the residue was dissolved in MeOH (1 mL) and
 15 purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to afford pure *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylacetamide (35 mg, yield: 80.3%). LC/MS (ESI⁺) 513.4 (M+H)⁺, *t*_R = 6.08
 20 min. HRMS C₂₇H₂₈O₃F₃N₄ (M+H)⁺ 513.2120 calcd for 513.2113.
¹H NMR (acetone-*d*₆) δ 7.49 (d, *J* = 9.1 Hz, 2H), 7.33 (m, 4H), 6.97 (d, *J* = 9.0 Hz, 2H), 4.15 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.58 (s, 1H), 3.49 (s, 1H), 3.16 (t, *J* = 6.6 Hz, 2H), 2.91, 2.80 (2 x s, 3H), 1.89, 1.50 (2 x s, 3H),
 25 0.87 (m, 2H), 0.78 (m, 2H) ppm.

Example 22

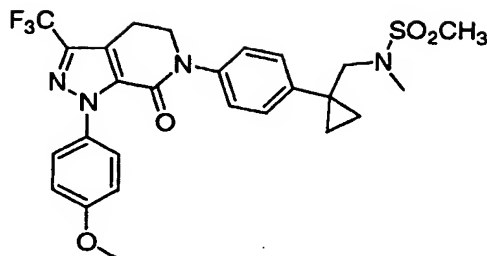
***N'*-ethyl-*N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylurea**



5 The product of Example 1 (20 mg, 0.042 mmol) was stirred in CH_2Cl_2 (1 mL) in a capped vial at RT. Et_3N (4 drops) was added followed by addition of ethyl isocyanide (2 drops). The resulting mixture was stirred at RT for 2 h. LC-MS
10 showed completion of the reaction. After evaporation of the solvents, the residue was dissolved in MeOH (1 mL), and purified by prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run) to afford pure *N'*-ethyl-*N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylurea (16
15 mg, yield: 70%). HRMS $\text{C}_{28}\text{H}_{31}\text{O}_3\text{F}_3\text{N}_5$ 542.2370 ($\text{M}+\text{H}$), calcd for 542.2380. ^1H NMR (acetone- d_6) δ 7.49 (d, $J = 9.2$ Hz, 2H), 7.30 (AA'BB', $J = 8.4$ Hz, 4H), 6.97 (d, $J = 8.8$ Hz, 2H), 4.14 (t, $J = 6.3$ Hz, 2H), 3.82 (s, 3H), 3.51 (s, 2H),
20 3.16 (t, $J = 6.4$ Hz, 2H), 3.02 (q, $J = 7.0$ Hz, 2H), 2.70 (s, 3H), 0.92 (t, $J = 7.0$ Hz, 3H), 0.86 (m, 2H), 0.76 (m, 2H) ppm.

Example 23

25 ***N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylethanesulfonamide**

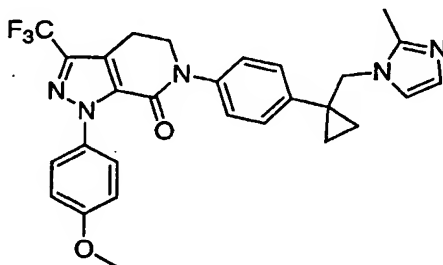


The product of Example 1 (20 mg, 0.042 mmol) was stirred in CH_2Cl_2 (1 mL) in a capped vial at RT. Pyridine (4 drops) was added followed by two drops of methanesulfonyl chloride. The resulting mixture was stirred for 20 min. LC-MS showed completion of the reaction. After evaporation of the solvents, the residue was dissolved in MeOH (1 mL), and purified by prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run) to afford pure *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylmethanesulfonamide (16 mg, yield: 69%). LC/MS (ESI⁺) 549.4 (M+H)⁺, t_R = 6.40 min.

15

Example 24

1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Part A. The product of part E in Example 1 (0.45 g, 0.98 mmol) was stirred in CH_2Cl_2 (10 mL) at 0°C under N_2 . PPh_3 (0.52 g, 2.0 eq) was added, followed by the addition of CBr_4 (0.33 g, 1.0 eq). The resulting mixture was stirred at 0°C for 30 min. LC-MS showed completion of the reaction. The mixture was extracted with EtOAc. The

organic layer was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness. It was used directly in the next step without purification.

LC/MS(ESI⁺) 520.4, 522.4 (M+H)⁺.

5

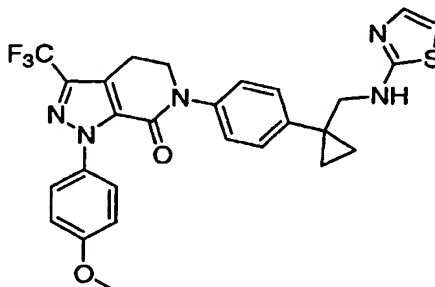
Part B. The product of Part A (0.20 g, 0.38 mmol), 2-methylimidazole (0.10 g, 1.22 mmol), and K₂CO₃ (0.25 g, 3.62 mmol) were stirred in DMF (0.4 mL) at RT under N₂. The mixture was heated at 85-90°C for 30 min. LC-MS showed completion of the reaction. After cooling to RT, H₂O was added. The mixture was purified by prep LC-MS (35-98% CH₃CN in H₂O) to give pure 1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one (53 mg, yield: 27%). LC/MS(ESI⁺) 522.4 (M+H)⁺. ¹H NMR (acetone-d₆) δ 7.53 (m, 1H), 7.49 (d, J = 8.8 Hz, 2H), 7.39 (m, 1H), 7.25 (AA'BB', J = 8.4 Hz, 4H), 6.98 (d, J = 8.8 Hz, 2H), 4.38 (s, 2H), 4.15 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.14 (t, J = 6.6 Hz, 2H), 2.10 (s, 3H), 1.26 (t, J = 5.5 Hz, 2H), 1.02 (t, J = 5.5 Hz, 2H) ppm.

20

Example 25

1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one

25

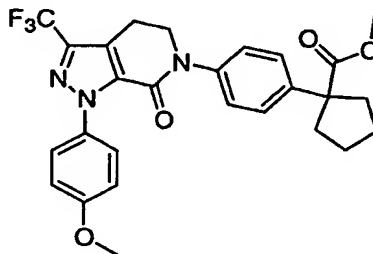


Following a procedure analogous to that of Example 24, the title compound was prepared by using 2-aminothiazole. The

product was purified by RP-prep LC-MS (35-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 540.6 (M+H)⁺, t_R = 3.37 min.

Example 26

5 methyl 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopentanecarboxylate



Part A. 1-Phenyl-cyclopentylcarboxylic acid (3.0 g, 15.8 mmol) was stirred in HOAc (10 mL) at RT under N₂. I₂ (4.01 g, 15.8 mmol) was added followed by the addition of NaIO₃ (0.78 g, 3.94 mmol) and conc. H₂SO₄ (0.3 mL). The resulting mixture was stirred at 70°C for 3 days. The cooled mixture was poured into H₂O, and extracted with EtOAc. The organic layer was washed with sodium thiosulfate and brine, dried over MgSO₄, filtered, and concentrated to dryness to yield 4-iodophenylcyclopentylcarboxylic acid (4.45 g, yield: 89%). LC/MS (ESI⁺) 317.6 (M+H)⁺.

Part B. The product from part A (1.08 g, 3.43 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.82 g, 2.64 mmol) were stirred in DMSO (3 mL) under N₂. K₂CO₃ (1.09 g, 7.90 mmol, 3.0 eq) was added followed by the addition of 1,10-phenanthroline (96 mg, 20 mol%) and CuI (100 mg, 20mol%). The resulting mixture was stirred at 130°C for 5h. LC-MS showed completion of the reaction. It was acidified with 1N HCl, and extracted with EtOAc (2x). The organic layer was washed with H₂O, brine, dried over MgSO₄, filtered, and concentrated to afford 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-

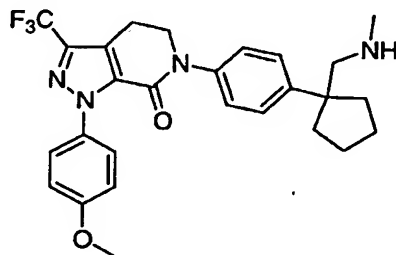
(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopentanecarboxylic acid (1.30 g, yield: 99%). LC/MS(ESI⁺) 500.6 (M+H)⁺.

5 Part C. The product of part B (40 mg, 0.080 mmol) was dissolved in MeOH (5 mL), and conc. HCl (0.5 mL) was added. The resulting mixture was stirred at 60°C overnight. After cooling, the mixture was purified by prep LC-MS (35-98% CH₃CN in H₂O in a 10-min run) to afford methyl
 10 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopentane carboxylate (32 mg, yield: 78%).
 LC/MS(ESI⁺) 514.6 (M+H)⁺, t_R = 6.09 min. ¹H NMR (CDCl₃) δ
 7.45 (d, J = 8.8 Hz, 2H), 7.30 (AA'BB', J = 8.6 Hz, 4H),
 15 6.92 (d, J = 9.0 Hz, 2H), 4.13 (t, J = 6.8 Hz, 2H), 3.81 (s, 3H), 3.39 (s, 3H), 3.15 (t, J = 6.6 Hz, 2H), 2.65-2.58 (m, 2H), 1.89-1.82 (m, 2H), 1.73-1.69 (m, 4H), 1.58 (m, 2H) ppm.

20

Example 27

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopentyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



25

Part A. The product from part B of Example 26 (1.46 g, 2.93 mmol) was stirred in THF (10 mL) at 0°C under N₂. Et₃N (0.62 mL, 4.40 mmol, 1.5 eq) was added followed by dropwise addition of ClCO₂Et (0.31 mL, 3.24 mmol, 1.1 eq). The
 30 reaction mixture was then stirred at 0°C for 1 h. TLC

showed completion of the reaction. The mixture was filtered, and rinsed with anhydrous THF. The THF filtrate (ca. 20 mL) was stirred at 0°C under N₂. MeOH (5 mL) was added followed by the addition of NaBH₄ (1.03 g, 27.10 mmol, 9.3 eq). The resulting mixture was stirred at 0°C for 40 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give 6-{4-[1-(hydroxymethyl)cyclopentyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.38 g, 97%). LC/MS (ESI⁺) 486.4 (M+H)⁺.

Part B. The product from part A (0.80 g, 1.65 mmol) was stirred in anhydrous CH₂Cl₂ (10 mL) at RT under N₂. NaOAc (0.5 g, 6.10 mmol) and molecular sieves (4Å, 1.2 g) were added followed by the addition of PCC (0.89 g, 4.12 mmol). The resulting slurry was stirred at RT for 4 h. Analytical LC-MS showed completion of the reaction. The mixture was filtered, and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to afford 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopentanecarbaldehyde (0.78 g, yield: 99%). LC/MS (ESI⁺) 484.6 (M+H)⁺.

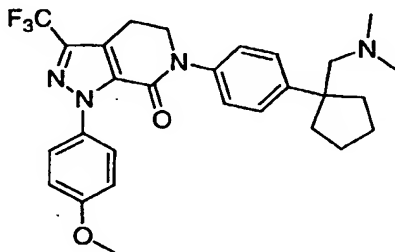
Part C. The product from part B (100 mg, 0.21 mmol) and methylamine hydrochloride (100 mg, excess) were stirred in dichloroethane (1.0 mL) in a capped vial. NaBH(OAc)₃ (200 mg, 0.94 mmol) was added followed by addition of three drops of HOAc. The reaction mixture was stirred at RT for 2h. Analytical LC-MS showed completion of the reaction. The mixture was evaporated and dissolved in aqueous MeOH.

It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to obtain 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopentyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (35 mg, yield: 34%). LC/MS (ESI⁺) 499.4 (M+H)⁺, t_R = 4.85 min. ¹H NMR (acetone-d₆) δ 7.50 (m, 4H), 7.35 (d, J = 8.4 Hz, 2H), 6.99 (d, J = 9.1 Hz, 2H), 4.17 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.39 (s, 2H), 3.16 (t, J = 6.3 Hz, 2H), 2.64 (s, 3H), 2.14-1.66 (m, 8H) ppm.

10

Example 28

6-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



15

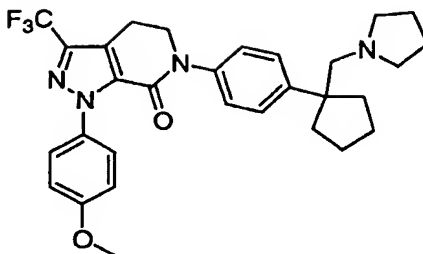
Following a procedure analogous to that used for Example 27, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 513.6 (M+H)⁺, t_R = 4.96 min. ¹H NMR (acetone-d₆) δ 7.52 (m, 4H), 7.38 (d, J = 8.5 Hz, 2H), 6.98 (d, J = 8.8 Hz, 2H), 4.19 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.60 (s, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.62 (s, 6H), 2.16-1.64 (m, 8H) ppm.

20

25

Example 29

1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopentyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

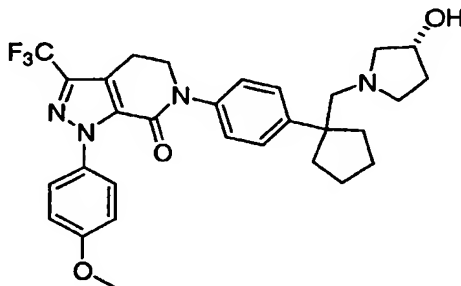


Following a procedure analogous to that used for Example 27, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

- 5 LC/MS (ESI⁺) 539.6 (M+H)⁺, *t_R* = 5.13 min. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.40 (d, *J* = 8.1 Hz, 2H), 6.99 (d, *J* = 8.8 Hz, 2H), 4.19 (t, *J* = 6.3 Hz, 2H), 3.83 (s, 3H), 3.75 (s, 2H), 3.52 (m, 2H), 3.17 (t, *J* = 6.3 Hz, 2H), 2.92 (m, 2H), 2.18 (m, 2H), 2.04-1.62 (m, 10H) ppm. ¹⁹F NMR
10 (acetone-*d*₆) δ -62.17 (TFA salt), -79.82 (CF₃) ppm.

Example 30

- 6-[4-(1-((3*R*)-3-hydroxy-1-pyrrolidinyl)methyl)cyclopentyl]phenyl]-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one, trifluoroacetic acid salt
- 15

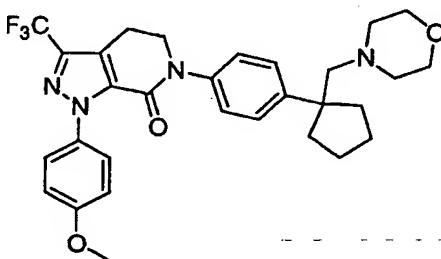


- Following a procedure analogous to that used for Example 27, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).
- 20 LC/MS (ESI⁺) 555.6 (M+H)⁺, *t_R* = 4.77 min. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.40 (d, *J* = 8.4 Hz, 2H), 6.99 (d, *J* = 8.8 Hz, 2H), 4.36 (s, br, 1H), 4.19 (t, *J* = 6.3 Hz, 2H), 3.83 (s, 3H), 3.72 (m, 2H), 3.59 (m, 2H), 3.17 (t, *J* = 6.3

Hz, 2H), 2.92 (m, 2H), 2.16-1.63 (m, 10H) ppm. ^{19}F NMR (acetone- d_6) δ -62.16 (TFA salt), -76.70 (CF_3) ppm.

Example 31

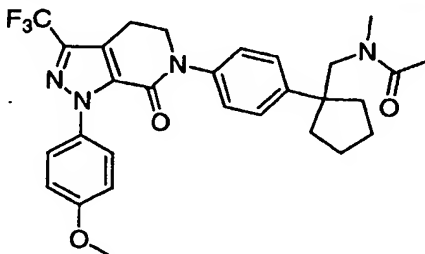
5 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopentyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



10 Following a procedure analogous to that used for Example 27, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run). LC/MS (ESI $^+$) 555.6 (M+H) $^+$, t_R = 4.49 min. ^1H NMR (acetone- d_6) δ 7.43 (m, 4H), 7.28 (d, J = 8.0 Hz, 2H), 6.89 (d, J = 8.4 Hz, 2H), 4.09 (t, J = 6.6 Hz, 2H), 3.74 (s, 3H), 3.65
15 (m, 4H), 3.50 (s, 2H), 3.33 (m, 2H), 3.07 (t, J = 6.3 Hz, 2H), 2.89 (m, 2H), 2.02-1.52 (m, 8H) ppm.

Example 32

20 N-[(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopentyl)methyl]-N-methylacetamide

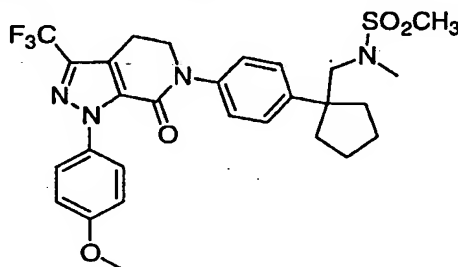


Following a procedure analogous to that used for Example 21, the title compound was prepared. The product was

purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).
 LC/MS (ESI⁺) 541.6 (M+H)⁺, t_R = 6.50 min. ¹H NMR (acetone-
 d₆) δ 7.49 (dd, J = 8.8, 1.8 Hz, 2H), 7.34 (m, 4H), 6.97
 (dd, J = 8.8, 1.8 Hz, 2H), 4.17 (t, J = 6 Hz, 2H), 3.82 (s,
 5 3H), 3.52 (s, 2H), 3.17 (t, J = 6 Hz, 2H), 2.34 (s, 3H),
 2.03-1.59 (m, 11H) ppm.

Example 33

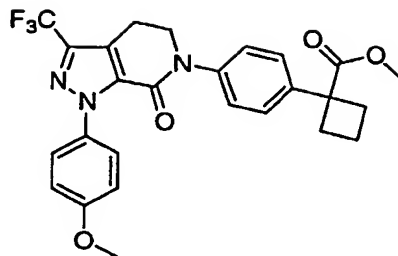
10 **N-[(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
 1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
 yl]phenyl)cyclopentyl)methyl]-N-methylmethanesulfonamide**



Following a procedure analogous to that used for Example
 23, the title compound was prepared. The product was
 15 purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).
 LC/MS (ESI⁺) 577.4 (M+H)⁺, t_R = 6.74 min. ¹H NMR (acetone-
 d₆) δ 7.50 (d, J = 9 Hz, 2H), 7.35 (m, 4H), 6.98 (d, J = 9
 Hz, 2H), 4.18 (t, J = 6.5 Hz, 2H), 3.83 (s, 3H), 3.22 (s,
 2H), 3.17 (t, J = 6.5 Hz, 2H), 2.70 (s, 3H), 2.17 (s, 3H),
 20 2.13 (m, 2H), 1.80 (m, 4H), 1.64 (m, 2H) ppm.

Example 34

**methyl 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
 1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
 25 yl]phenyl)cyclobutanecarboxylate**



Part A. 1-Phenyl-1-cyclobutylcarbonitrile (5.0 g, 31.83 mmol) and KOH (85%, 6.29 g, 95.49 mmol, 3 eq) were heated in ethylene glycol (10 mL) at 185-190°C for 6h under N₂.

5 LC-MS showed completion of the reaction. H₂O was added to the cooled mixture. It was extracted with Et₂O (3x). The aqueous layer was acidified with conc. HCl, and then extracted with CHCl₃ (2x). The chloroform layer was washed with H₂O, brine, dried over MgSO₄, filtered, and
10 concentrated to dryness to give 1-phenyl-1-cyclobutyl carboxylic acid (4.43 g, yield: 79.2%). LC/MS (ESI⁺) 177.4 (M+H)⁺, t_R = 2.56 min (10-90% CH₃CN/H₂O in a 6-min run).

Part B. The product from part A (4.43 g, 25.2 mmol) was
15 stirred in HOAc (20 mL) at RT under N₂. I₂ (6.40 g, 25.2 mmol) was added, followed by the addition of NaIO₃ (1.25 g, 6.3 mmol) and conc. H₂SO₄ (0.5 mL). The resulting mixture was stirred at 70°C for 2 days. LC-MS showed completion of the reaction. The cooled mixture was poured into H₂O, and
20 extracted with EtOAc. The organic layer was washed with sodium thiosulfate, brine, dried over MgSO₄, filtered, and concentrated to dryness to give 4-iodophenylcyclobutyl carboxylic acid (6.49 g, 85%). LC/MS (ESI⁺) 303.2 (M+H)⁺, t_R = 2.55 min (10-90% CH₃CN/H₂O in a 4-min run).

25

Part C. The product from part B (1.20 g, 3.97 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.87 g, 2.8 mmol) were stirred in DMSO (3 mL) under N₂. K₂CO₃ (1.16 g, mmol, 3.0

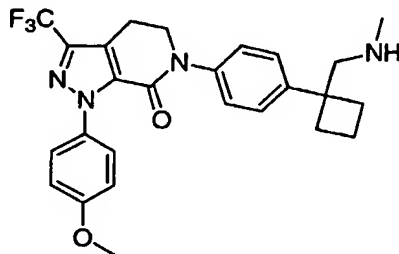
eq) was added followed by the addition of 1,10-phenanthroline (100 mg, 20 mol%) and CuI (106 mg, 20 mol%). The resulting mixture was stirred at 130°C overnight. LC-MS showed completion of the reaction. EtOAc was added to the cooled solution. The solution was acidified with 1N HCl, and the organic layer was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to afford 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclobutane carboxylic acid (1.34 g, yield 97%). LC/MS (ESI⁺) 486.6 (M+H)⁺, t_R = 2.81 min (10-90% CH₃CN/H₂O in a 4-min run).

Part D. The product from part C (50 mg, 0.103 mmol) was dissolved in MeOH (5 mL), and conc. HCl (0.5 mL) was added. The resulting mixture was stirred at reflux for 2 h. After cooling, the mixture was purified by prep LC-MS (35-98% CH₃CN in H₂O) to afford the title compound (35 mg, yield: 68%). LC/MS (ESI⁺) 499.4 (M+H)⁺, t_R = 5.70 min. ¹H NMR (CDCl₃) δ 7.45 (d, J = 8.8 Hz, 2H), 7.28 (AA'BB', J = 8Hz, 4H), 6.91 (d, J = 8.8 Hz, 2H), 4.13 (t, J = 6.9 Hz, 2H), 3.80 (s, 3H), 3.63 (s, 3H), 3.15 (t, J = 6.6 Hz, 2H), 2.81 (m, 2H), 2.48 (m, 2H), 2.03 (m, 1H), 1.86 (m, 1H) ppm.

25

Example 35

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclobutyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



30

Part A. The product from part C of Example 34 (1.32 g, 2.72 mmol) was stirred in THF (10 mL) at 0°C under N₂. Et₃N (0.59 mL, 4.08 mmol, 1.5 eq) was added followed by dropwise addition of ClCO₂Et (0.38 mL, 3.54 mmol, 1.3 eq). The
5 reaction mixture was then stirred at 0°C for 30 min. TLC showed completion of the reaction. The mixture was filtered and rinsed with anhydrous THF. The THF filtrate (ca. 20 mL) was stirred at 0°C under N₂. MeOH (4 mL) was added followed by the addition of NaBH₄ (1.03 g, 27.10
10 mmol, 10 eq). The resulting mixture was stirred at 0°C for 40 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered,
15 and concentrated to dryness to give 6-{4-[1-(hydroxymethyl)cyclobutyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.30 g, 99%). LC/MS (ESI⁺) 472.6 (M+H)⁺, t_R = 2.84 min (10-90% CH₃CN/H₂O in a 4-min run).

20

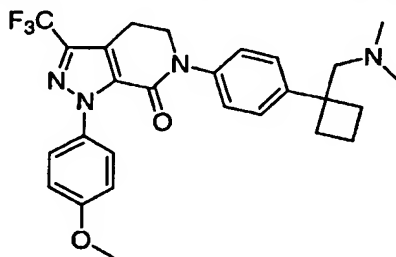
Part B. The product from part A (0.90 g, 1.91 mmol) was stirred in anhydrous CH₂Cl₂ (10 mL) at RT under N₂. NaOAc (0.32 g, 3.82 mmol, 2.0 eq) and molecular sieves (4A, 0.90 g) were added followed by the addition of PCC (0.69 g, 2.87
25 mmol, 1.5 eq). The resulting slurry was stirred at RT for 4 h. Analytical LC-MS showed completion of the reaction. The mixture was filtered and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to afford
30 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclobutanecarbaldehyde (0.88 g, yield: 99%). LC/MS (ESI⁺) 470.6 (M+H)⁺, t_R = 3.01 min (10-90% CH₃CN/H₂O in a 4-min run).

Part C. The product from part B (500 mg, 1.04 mmol), methylamine hydrochloride (200 mg, excess) were stirred in dichloroethane (15 mL) at RT under N₂. NaBH(OAc)₃ (1.03 g, 4.86 mmol) was added followed by addition of three drops of HOAc. The reaction mixture was stirred at RT for 2.5 h. Analytical LC-MS showed completion of the reaction. The mixture was evaporated, and dissolved in aqueous MeOH. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to obtain 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclobutyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (230 mg, 46%). LC/MS (ESI⁺) 485.4 (M+H)⁺, t_R = 4.93 min. ¹H NMR (acetone-d₆) δ 7.51 (d, 2H), 7.33 (m, 4H), 6.99 (d, 2H), 4.17 (m, 2H), 3.83 (s, 3H), 3.62 (m, 2H), 3.17 (m, 2H), 2.73 (s, 3H), 2.46 (m, 4H), 2.15-1.86 (m, 2H) ppm. ¹⁹F NMR (acetone-d₆) δ -62.18 (TFA), -76.65 (CF₃) ppm.

20

Example 36

6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



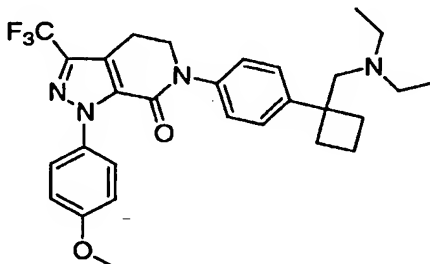
25 Following a procedure analogous to that used for Example 35, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 499.6 (M+H)⁺, t_R = 4.75 min. ¹H NMR (acetone-d₆) δ 7.49 (m, 4H), 7.39 (d, J = 8.0 Hz, 2H), 6.98 (d, J =

9.1 Hz, 2H), 4.19 (t, J = 6.4 Hz, 2H), 3.83 (s, 3H), 3.77 (m, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.68 (s, 6H), 2.48 (t, J = 7.5 Hz, 4H), 2.09 (m, 1H), 1.89 (m, 1H) ppm.

5

Example 37

6-(4-{1-[(diethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

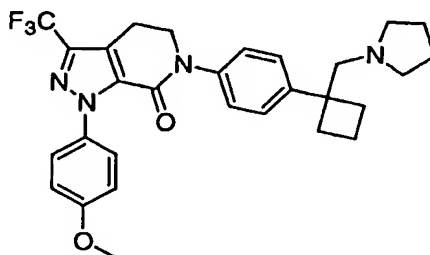


10 Following a procedure analogous to that used for Example 35, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 527.6 (M+H)⁺, t_R = 5.04 min. ¹H NMR (acetone-*d*₆) δ 7.53 (m, 4H), 7.39 (d, J = 8.5 Hz, 4H), 6.98 (d, J = 9.1 Hz, 2H), 4.19 (t, J = 6.3 Hz, 2H), 3.83 (s, 3H), 3.68 (s, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.95 (m, 4H), 2.48 (t, J = 7.8 Hz, 4H), 2.10-1.85 (m, 2H), 1.16 (m, 6H) ppm.

20

Example 38

1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclobutyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



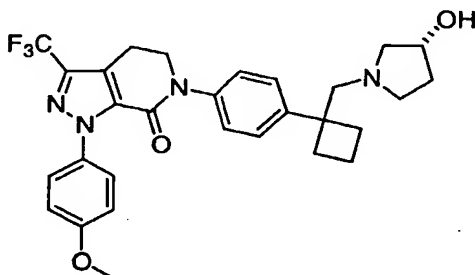
Following a procedure analogous to that used for Example 35, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS (ESI⁺) 525.6 (M+H)⁺, t_R = 4.97 min. ¹H NMR (acetone-
 5 d₆) δ 7.51 (d, J = 8.4 Hz, 2H), 7.39 (m, 4H), 6.98 (d, J = 8.8 Hz, 2H), 4.19 (t, J = 6.4 Hz, 2H), 3.87 (m, 2H), 3.83 (s, 3H), 3.51 (m, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.93 (m, 2H), 2.47 (m, 4H), 2.09-1.85 (m, 6H) ppm. ¹⁹F NMR (acetone-
 d₆) δ -62.16 (TFA), -76.74 (CF₃) ppm.

10

Example 39

6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclobutyl)phenyl]-1-(4-methoxyphenyl)-
 3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
 15 c]pyridin-7-one, trifluoroacetic acid salt



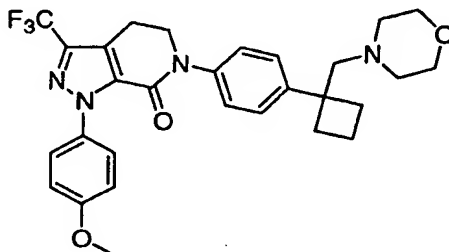
Following a procedure analogous to that used for Example 35, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS (ESI⁺) 541.6 (M+H)⁺, t_R = 4.77 min. ¹H NMR (acetone-
 20 d₆) δ 7.51 (d, J = 8.9 Hz, 2H), 7.39 (m, 4H), 6.98 (d, J = 9.1 Hz, 2H), 4.33 (m, 1H), 4.20 (t, J = 6.5 Hz, 2H), 3.83 (m, 7H), 3.52 (m, 2H), 3.17 (t, J = 6.5 Hz, 2H), 2.47 (m, 4H), 2.14-1.84 (m, 4H). ¹⁹F NMR (acetone-d₆) δ -62.16

25 (TFA), -76.34 (CF₃) ppm.

Example 40

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

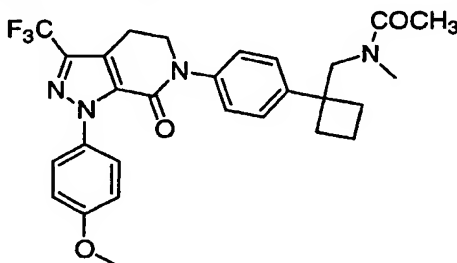


Following a procedure analogous to that used for Example 35, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

LC/MS (ESI⁺) 541.6 (M+H)⁺, t_R = 4.86 min. ¹H NMR (acetone-d₆) δ 7.49 (m, 4H), 7.37 (m, 2H), 6.98 (d, J = 8.8 Hz, 2H), 4.18 (m, 2H), 3.83 (s, 3H), 3.74 (m, 8H), 3.17 (t, J = 6.5 Hz, 2H), 3.00 (m, 2H), 2.46 (m, 4H), 1.86 (m, 2H) ppm.

Example 41

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclobutyl)methyl]-N-methylacetamide

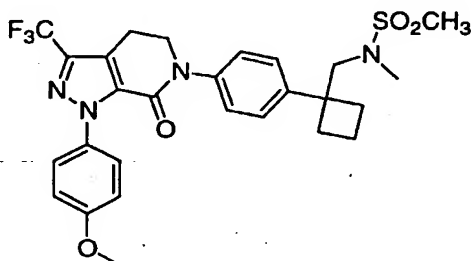


Following a procedure analogous to that used for Example 21, the title compound was prepared by using the product from Example 37 and acetyl chloride as the starting material. The mixture was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 527.2 (M+H)⁺, t_R = 6.36 min. ¹H NMR (acetone-d₆) δ 7.50 (d, J = 9.2 Hz, 2H),

7.33 (d, $J = 8.7$ Hz, 2H), 7.24 (m, 2H), 6.98 (d, $J = 9.2$ Hz, 2H), 4.17 (d, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.71, 3.67 (s, 2H), 3.17 (t, $J = 6.5$ Hz, 2H), 2.76, 2.45 (s, 3H), 2.35 (m, 2H), 2.18 (m, 1H), 2.08 (m, 1H), 1.76 (m, 2H), 1.33 (m, 1H) ppm.

Example 42

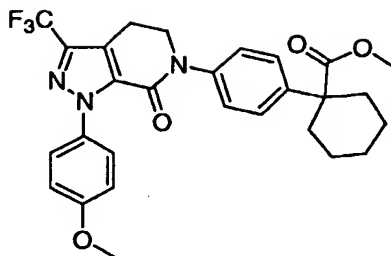
***N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclobutyl)methyl]-*N*-methylmethanesulfonamide**



Following a procedure analogous to that used for Example 23, the title compound was prepared. The mixture was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 563.2 (M+H)⁺, $t_R = 6.62$ min. ¹H NMR (acetone-*d*₆) δ 7.50 (d, $J = 8.8$ Hz, 4H), 7.34 (d, $J = 8.5$ Hz, 2H), 7.23 (d, $J = 8.4$ Hz, 2H), 6.98 (d, $J = 8.8$ Hz, 2H), 4.18 (d, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.45 (s, 2H), 3.17 (t, $J = 6.5$ Hz, 2H), 2.70 (s, 3H), 2.38 (m, 2H), 2.28 (s, 3H), 2.26 (m, 2H), 2.03 (m, 1H), 1.81 (m, 1H) ppm.

Example 43

1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]-phenyl}cyclohexanecarboxylic acid methyl ester



Part A. 1-Phenyl-cyclohexylcarboxylic acid (3.0 g, 14.70 mmol) was stirred in HOAc (10 mL) at RT under N₂. I₂ (3.73 g, 14.70 mmol) was added, followed by the addition of NaIO₃ (0.72 g, 3.64 mmol) and conc. H₂SO₄ (0.2 mL). The resulting mixture was stirred at 70°C for 2 days. LC-MS showed the majority was the desired product. After partial evaporation, the cooled mixture was poured into H₂O and extracted with EtOAc. The organic layer was washed with sodium thiosulfate and brine, dried over MgSO₄, filtered, and concentrated to dryness to give almost pure 4-iodophenylcyclohexylcaroxylic acid (4.56 g, yield: 93.7%). LC/MS (ESI⁺) 331.4 (M+H)⁺, t_R = 3.96 min (10-90% CH₃CN/H₂O in a 6-min run).

15

Part B. The product of part A (0.70 g, 2.25 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.70 g, 2.25 mmol) were stirred in DMSO (3 mL) under N₂. K₂CO₃ (0.93 g, mmol, 3.0 eq) was added, followed by the addition of 1,10-phenanthroline (80 mg, 20 mmol%) and CuI (85 mg, 20mmol%). The resulting mixture was stirred at 130°C for 2 days. LC-MS showed completion of the reaction. EtOAc was added to the cooled solution. It was acidified with 1N HCl; and the organic layer was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated. LC/MS (ESI⁺) 514.4 (M+H)⁺. The residue (50 mg) was dissolved in MeOH (5 mL), and conc. HCl (0.5 mL) was added. The resulting mixture was stirred at 60°C for 4 h. After cooling, the mixture was purified by prep LC-MS (35-98% CH₃CN in H₂O) to give

30

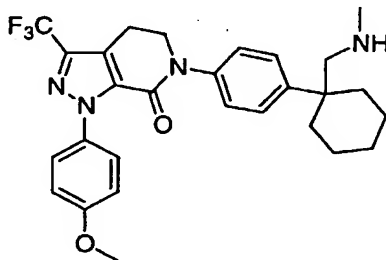
pure 1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}cyclohexanecarboxylic acid methyl ester (43 mg, yield: 83.7%). LC/MS (ESI⁺) 528.4 (M+H)⁺, t_R = 6.38 min.

5 ¹H NMR (CDCl₃) δ 7.45 (d, J = 9.1 Hz, 2H), 7.32 (AA'BB', J = 8.6 Hz, 4H), 6.92 (d, J = 8.8 Hz, 2H), 4.13 (t, J = 6.6 Hz, 2H), 3.81 (s, 3H), 3.63 (s, 3H), 3.15 (t, J = 6.6 Hz, 2H), 2.45 (m, 2H), 1.72-1.24 (m, 8H) ppm.

10

Example 44

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclohexyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



15

Part A. The product from part B of Example 43 (1.34 g, 2.61 mmol) was stirred in THF (10 mL) at 0°C under N₂. Et₃N (0.55 mL, 3.92 mmol, 1.5 eq) was added followed by dropwise addition of ClCO₂Et (0.33 mL, 3.34 mmol, 1.3 eq). The

20 reaction mixture was then stirred at 0°C for 20 min. TLC showed completion of the reaction. The mixture was filtered, and rinsed with anhydrous THF. The THF filtrate (ca. 20 mL) was stirred at 0°C under N₂. MeOH (3.5 mL) was added followed by the addition of NaBH₄ (1.00 g, 26.3 mmol, 25 10 eq). The resulting mixture was stirred at 0°C for 40 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to

dryness to give 6-{4-[1-(hydroxymethyl)cyclohexyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.21 g, yield 92.8%).

LC/MS (ESI⁺) 500.6 (M+H)⁺, t_R = 3.06 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (CDCl₃) δ 7.46 (d, J = 8.8 Hz, 2H), 7.34 (AA'BB', J = 8.4 Hz, 4H), 6.92 (d, J = 8.8 Hz, 2H), 4.15 (t, J = 6.7 Hz, 2H), 3.81 (s, 3H), 3.49 (s, 2H), 3.16 (t, J = 6.6 Hz, 2H), 2.24 (m, 2H), 2.13 (m, 2H), 1.56 (m, 4H), 1.34 (m, 2H) ppm.

10

Part B. The product from part A (0.56 g, 1.12 mmol) was stirred in anhydrous CH₂Cl₂ (10 mL) at RT under N₂. NaOAc (0.37 g, 4.48 mmol, 4 eq) and molecular sieves (4Å, 1.0 g) were added followed by the addition of PCC (0.73 g, 3.36 mmol, 3 eq). The resulting slurry was stirred at RT for 1.5h. Analytical LC-MS showed completion of the reaction. The mixture was filtered, and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to afford 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclohexanecarbaldehyde (0.54 g, yield: 100%). LC/MS (ESI⁺) 498.6 (M+H)⁺, t_R = 3.20 min (10-90% CH₃CN in H₂O in a 4-min run).

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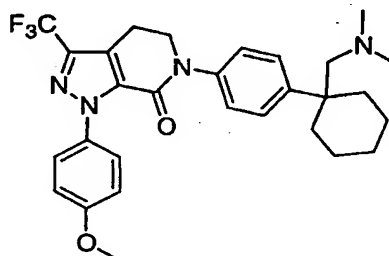
Part C. The product from part B (0.4 g, 0.85 mmol) and methylamine hydrochloride (0.2 mg, 2.99 mmol, excess) were stirred in dichloroethane (8 mL) at RT under N₂. NaBH(OAc)₃ (0.85 mg, 4.01 mmol) was added followed by addition of HOAc (0.1 mL). The reaction mixture was stirred at RT for 2h. Analytical LC-MS showed completion of the reaction. The mixture was evaporated, and dissolved in aqueous MeOH. The mixture was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to afford 1-(4-methoxyphenyl)-6-(4-{1-

30

[(methylamino)methyl]cyclohexyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (120 mg. Yield: 29%). LC/MS (ESI⁺) 513.4 (M+H)⁺, t_R = 4.97 min. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.38 (d, J = 8.5 Hz, 2H), 6.97 (d, J = 8.8 Hz, 2H), 4.19 (t, J = 6.3 Hz, 2H), 3.81 (s, 3H), 3.60 (s, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.62 (s, 6H), 2.16-1.64 (m, 8H) ppm.

Example 45

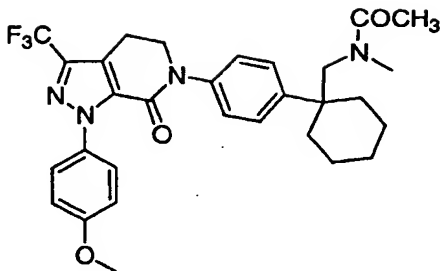
6-(4-{1-[(dimethylamino)methyl]cyclohexyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



Following a procedure analogous to that used for Example 44, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 526.4 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.38 (d, J = 8.5 Hz, 2H), 6.97 (d, J = 8.8 Hz, 2H), 4.19 (t, J = 6.3 Hz, 2H), 3.81 (s, 3H), 3.60 (s, 2H), 3.17 (t, J = 6.3 Hz, 2H), 2.62 (s, 6H), 2.16-1.64 (m, 8H) ppm.

Example 46

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclohexyl)methyl]-*N*-methylacetamide



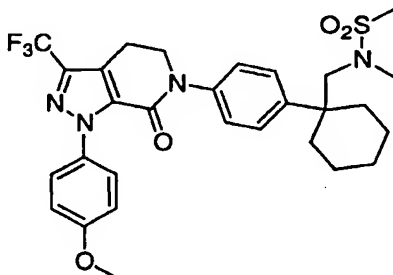
Following a procedure analogous to that used for Example 21, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

5 LC/MS (ESI⁺) 555.2 (M+H)⁺, *t_R* = 6.76 min. ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J* = 8.8 Hz, 2H), 7.38 (m, 4H), 6.98 (d, *J* = 9.2 Hz, 2H), 4.19 (t, *J* = 6.4 Hz, 2H), 3.83 (s, 3H), 3.39 (m, 2H), 3.17 (t, *J* = 6.4 Hz, 2H), 2.70, 2.42 (s, 3H), 2.02, 1.93 (m, 3H), 1.56 (m, 6H), 1.40 (m, 4H) ppm.

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Example 47

***N*-[(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclohexyl)methyl]-*N*-methylmethanesulfonamide**



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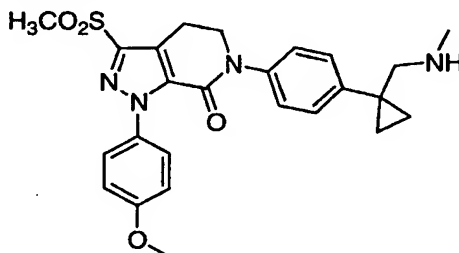
Following a procedure analogous to that used for Example 23, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

15 LC/MS (ESI⁺) 591.2 (M+H)⁺, *t_R* = 6.90 min. ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J* = 8.8 Hz, 2H), 7.42 (AA'BB', *J* = 8.8 Hz, 4H), 6.98 (d, *J* = 8.7 Hz, 2H), 4.19 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.17 (t, *J* = 6.6 Hz, 2H), 3.10 (s, 2H), 2.69 (m, 3H), 2.21 (m, 3H), 1.57 (m, 6H), 1.29 (m, 4H) ppm.

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Example 48

1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopropyl}phenyl)-3-
(methylsulfonyl)-1,4,5,6-tetrahydro 7H-pyrazolo[3,4-
c]pyridin-7-one, trifluoroacetic acid salt



Part A. 3-Chloro-5,6-dihydro-2(1H)-pyridinone (10.0 g, 38.17 mmol) and (1Z)-1-[chloro(methylsulfonyl)methylene]-2-(4-methoxyphenyl)hydrazine (5.0 g, 38.17 mmol) were stirred in toluene (200 mL) at RT under N₂. Et₃N (30 mL, 215.24 mmol) in toluene (150 mL) was added dropwise to the solution. After addition, the mixture was heated at 85°C overnight. After cooling, H₂O was added. It was extracted with EtOAc (2x). The organic layers were washed with H₂O (2x) and brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by flash column chromatography (silica gel, CH₂Cl₂, then CH₂Cl₂: EtOAc = 1:1, then EtOAc) to give 1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (3.85 g, yield: 31%). LC/MS (ESI⁺) 322.4 (M+H)⁺, t_R = 1.79 min (10-90% CH₃CN/H₂O in a 4-min run).

Part B. The product from part A (2.07 g, 6.23 mmol) and 4-iodophenylcyclopropylcarboxylic acid (2.75 g, 9.54 mmol, 1.5 eq) were stirred in DMSO (6 mL) under N₂. K₂CO₃ (2.57 g, 18.62 mmol, 3.0 eq) was added, followed by the addition of CuI (0.24 g, 20mol%) and 1,10-phenanthroline (0.23 g, 20 mol%). The resulting mixture was heated at 130°C overnight. After cooling, 1N HCl was added to acidify the solution. It was extracted with EtOAc (2x), washed with H₂O and

brine, dried over MgSO_4 , filtered, and concentrated to dryness to give 1-(4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropanecarboxylic acid (2.95 g, 5 yield: 95%). LC/MS (ESI^+) 482.4 ($\text{M}+\text{H}^+$), $t_R = 2.25$ min (10-90% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 4-min run).

Part C. The product from part B (2.21 g, 4.59 mmol) was stirred in THF (15 mL) at 0°C under N_2 . Et_3N (0.96 mL, 6.89 10 mmol, 1.5 eq) was added, followed by dropwise addition of ClCO_2Et (0.57 mL, 5.48 mmol, 1.2 eq). The reaction mixture was then stirred at 0°C for 40 min. TLC showed completion of the reaction. The mixture was filtered, and rinsed with anhydrous THF. The THF filtrate (ca. 20 mL) was stirred at 15 0°C under N_2 . MeOH (4 mL) was added, followed by portionwise addition of NaBH_4 (1.62 g, 42.63 mmol, 9 eq). The resulting mixture was stirred at 0°C for 35 min. Analytical LC-MS showed completion of the reaction. Sat'd Na_2SO_4 was then added. The mixture was extracted with EtOAc 20 (2x). The organic layer was washed with H_2O (2x) and brine (2x), dried over Na_2SO_4 , filtered, and concentrated to dryness to give 6-(4-[1-(hydroxymethyl)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.23 g, 84.7%). LC/MS 25 (ESI^+) 468.4 ($\text{M}+\text{H}^+$), $t_R = 2.24$ min (10-90% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 4-min run).

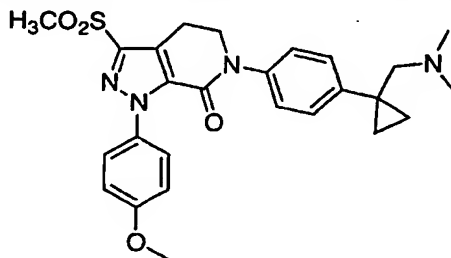
Part D. The product from part C (1.52 g, 3.27 mmol) was stirred in anhydrous CH_2Cl_2 (15 mL) at RT under N_2 . NaOAc 30 (0.54 g, 6.54 mmol, 2.0 eq) and molecular sieves (4Å, 1.5 g) were added, followed by the addition of PCC (1.06 g, 4.90 mmol, 1.5 eq). The resulting slurry was stirred at RT for 2 h. Analytical LC-MS showed completion of the reaction. The mixture was filtered through Celite®, and

rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give 1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropanecarbaldehyde (1.35 g, yield: 89%). LC/MS (ESI⁺) 466.4 (M+H)⁺, t_R = 2.38 min (10-90% CH₃CN/H₂O in a 4-min run).

Part E. The product from Part E (100 mg, 0.22 mmol) and methylamine hydrochloride (50 mg, excess) were stirred in dichloroethane (1 mL) in a capped vial. NaBH(OAc)₃ (250 mg, 1.16 mmol) was added followed by addition of one drop of HOAc. The reaction mixture was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was evaporated, and dissolved in aqueous MeOH. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to obtain the product 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7H-pyrazolo[3,4-c]pyridin-7-one (33 mg, 31.3%). LC/MS (ESI⁺) 481.4 (M+H)⁺, t_R = 3.99 min. ¹H NMR (acetone-d₆) δ 7.53 (m, 4H), 7.32 (d, J = 7.7 Hz, 2H), 7.00 (d, J = 8.8 Hz, 2H), 4.13 (t, J = 6.4 Hz, 2H), 3.83 (s, 3H), 3.46 (m, 2H), 3.26 (m, 5H), 2.80 (m, 3H), 1.11 (m, 2H), 1.00 (m, 2H) ppm.

Example 49

6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



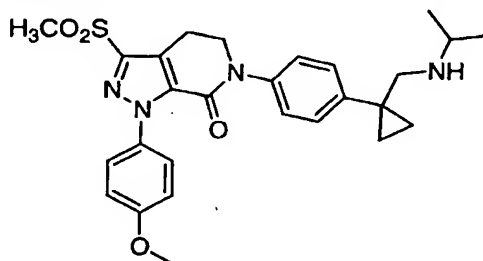
Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 495.6 (M+H)⁺, t_R = 4.06 min.

5 ¹H NMR (acetone-d₆) δ 7.55 (m, 4H), 7.38 (d, J = 7.5 Hz, 2H), 7.02 (d, J = 8.4 Hz, 2H), 4.16 (d, J = 6.6 Hz, 2H), 3.86 (m, 3H), 3.71 (m, 2H), 3.64 (m, 2H), 3.29, 3.25 (m, 6H), 3.09 (t, J = 6.6 Hz, 2H), 2.99 (m, 3H), 1.17 (m, 2H), 1.13 (m, 2H) ppm.

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Example 50

6-(4-(1-[(isopropylamino)methyl]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



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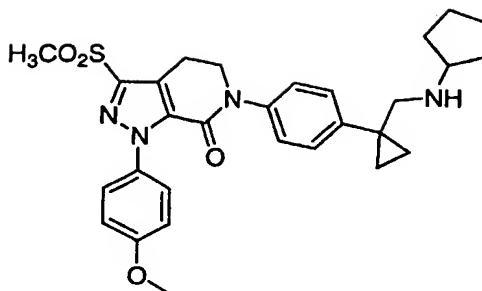
Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 509.6 (M+H)⁺, t_R = 4.34 min.

20 ¹H NMR (acetone-d₆) δ 7.58 (m, 4H), 7.33 (d, J = 8.4 Hz, 2H), 7.02 (d, J = 9.1 Hz, 2H), 4.52 (m, 1H), 4.16 (d, J = 6.6 Hz, 2H), 3.86 (s, 3H), 3.79 (m, 2H), 3.61 (m, 1H), 3.48 (m, 2H), 3.29 (m, 5H), 1.33 (d, J = 6.2 Hz, 2H), 1.17 (m, 2H), 1.07 (m, 2H) ppm.

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Example 51

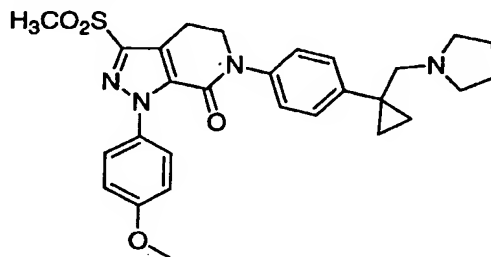
6-(4-(1-[(cyclopentylamino)methyl]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt,



Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 535.4 (M+H)⁺, *t_R* = 4.30 min. ¹H NMR (acetone-*d*₆) δ 7.55 (m, 4H), 7.33 (d, *J* = 7.4 Hz, 2H), 7.02 (d, *J* = 8.8 Hz, 2H), 4.16 (t, *J* = 6.6 Hz, 2H), 3.85 (s, 3H), 3.64 (m, 1H), 3.44 (m, 2H), 3.09 (m, 2H), 2.05 (m, 2H), 1.70 (m, 4H), 1.53 (m, 2H), 1.16 (m, 2H), 0.98 (m, 2H) ppm.

Example 52

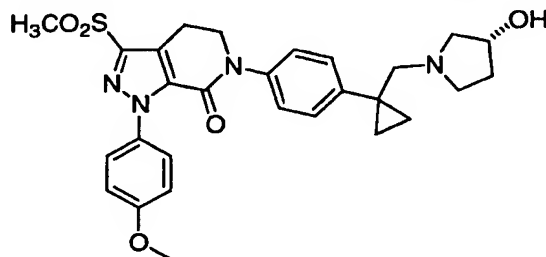
1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 521.4 (M+H)⁺, *t_R* = 4.07 min. ¹H NMR (acetone-*d*₆) δ 7.58 (m, 4H), 7.37 (d, *J* = 7.7 Hz, 2H), 7.02 (d, *J* = 7.7 Hz, 2H), 4.19 (t, *J* = 6.6 Hz, 2H), 3.85 (s, 3H), 3.70 (m, 4H), 3.28 (m, 5H), 3.09 (t, *J* = 6.6 Hz, 2H), 2.08-2.03 (m, 2H), 1.16 (m, 2H), 1.06 (m, 2H) ppm.

Example 53

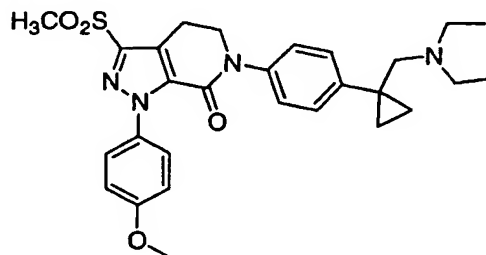
6-[4-(1-[[[(3R)-3-hydroxy-1-pyrrolidinyl]methyl]cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 537.6 (M+H)⁺, t_R = 3.90 min. ¹H NMR (acetone-d₆) δ 7.55 (m, 4H), 7.38 (m, 2H), 7.01 (d, J = 8.5 Hz, 2H), 4.52 (m, 1H), 4.17 (m, 2H), 3.85 (s, 3H), 3.79 (m, 2H), 3.63 (m, 2H), 3.29 (m, 5H), 3.09 (m, 2H), 2.17 (m, 2H), 1.17 (m, 2H), 1.07 (m, 2H) ppm.

Example 54

6-(4-(1-[(diethylamino)methyl]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



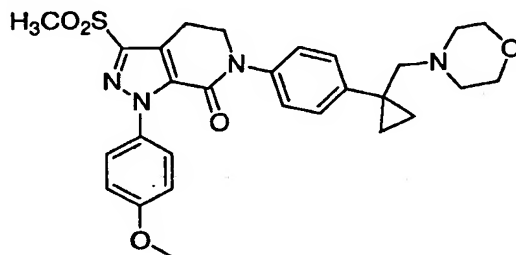
Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 523.4 (M+H)⁺, t_R = 4.54 min.

^1H NMR (acetone- d_6) δ 7.55 (m, 4H), 7.35 (d, J = 7.5 Hz, 2H), 7.00 (d, J = 8.8 Hz, 2H), 4.16 (d, J = 6.6 Hz, 2H), 3.84 (s, 3H), 3.56 (m, 4H), 3.28 (m, 3H), 3.10 (m, 2H), 1.17 (m, 5H), 1.17 (m, 2H), 1.05 (m, 2H) ppm.

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Example 55

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-(4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



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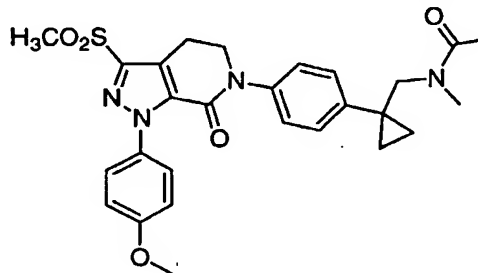
Following a procedure analogous to that used for the preparation of Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run). LC/MS (ESI $^+$) 537.6 (M+H) $^+$, t_R = 4.18 min.

^1H NMR (acetone- d_6) δ 7.54 (m, 4H), 7.36 (d, J = 7.7 Hz, 2H), 7.00 (d, J = 8.8 Hz, 2H), 4.17 (t, J = 6.6 Hz, 2H), 3.83 (m, 7H), 3.68 (m, 2H), 3.61 (m, 2H), 3.25 (m, 5H), 3.15 (m, 2H), 2.08-2.03 (m, 2H), 1.18 (m, 2H), 1.07 (m, 2H) ppm.

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Example 56

N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)methyl]-N-methylacetamide



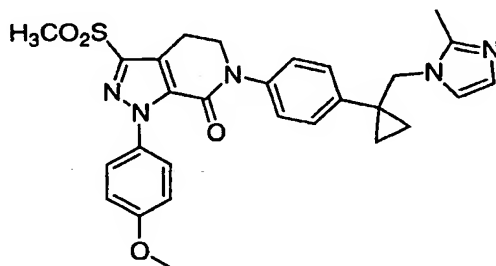
25

Following a procedure analogous to that used for the preparation of Example 21, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 523.6 (M+H)⁺, t_R = 5.07 min.

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Example 57

3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



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Part A. The product of part C in example 48 (0.69 g, 1.48 mmol) was stirred in CH₂Cl₂ (6 mL) at 0°C under N₂. PPh₃ (0.50 g, 1.91 mmol, 1.3 eq) was added, followed by the addition of CBr₄ (0.49 g, 1.48 mmol, 1.0 eq). The resulting mixture was stirred at 0°C for 30 min. LC-MS showed completion of the reaction (10-90% CH₃CN in H₂O in a 4-min run, t_R = 2.73 min). Sat'd NH₄Cl was then added. The mixture was extracted with EtOAc. The organic layer was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness (0.39 g, yield: 50%). The product was used directly in the next step. LC/MS (ESI⁺) 530.2, 532.2 (M+H)⁺.

Part B. The product of Part A (0.13 g, 0.25 mmol), 2-methylimidazole (50 mg, 0.64 mmol), and K₂CO₃ (0.13 g, 1.0 mmol) were stirred in DMF (0.4 mL) under N₂. The mixture was heated at 85-90°C for 30 min. LC-MS showed completion of the reaction. After cooling to RT, H₂O was added. The mixture was purified by prep LC-MS (15-70% CH₃CN in H₂O) to

give pure 3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one (36 mg, yield %).

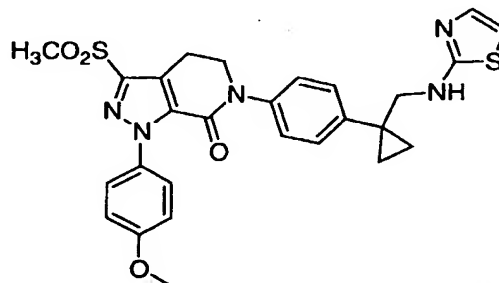
LC/MS (ESI⁺) 532.4 (M+H)⁺, *t_R* = 4.43 min. ¹H NMR (acetone-

5 *d*₆) δ 7.57 (m, 1H), 7.52 (d, *J* = 8.8 Hz, 2H), 7.45 (m, 1H), 7.26 (AA'BB', *J* = 8.4 Hz, 4H), 6.99 (d, *J* = 9.1 Hz, 2H), 4.41 (s, 2H), 4.13 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.25 (m, 5H), 2.11 (s, 3H), 1.27 (t, *J* = 5.8 Hz, 2H), 1.03 (t, *J* = 5.8 Hz, 2H) ppm.

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Example 58

3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



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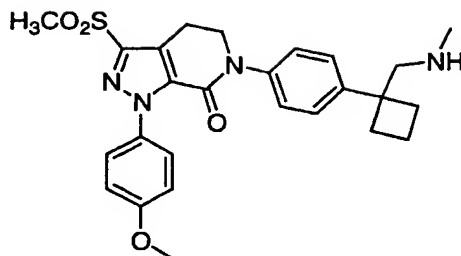
Following a procedure analogous to that of Example 25, the title compound was prepared. The product was purified by RP-prep LC-MS (35-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 550.4 (M+H)⁺, *t_R* = 2.36 min.

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Example 59

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclobutyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

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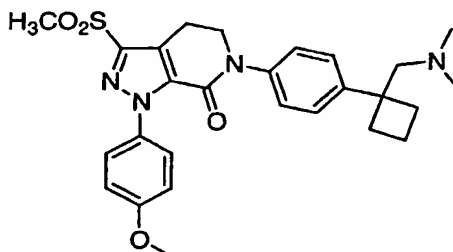


Following a procedure analogous to that used for Example 48, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 4.04 min). LC/MS (ESI⁺) 495.4 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.53 (d, J = 8.8 Hz, 4H), 7.32 (m, 4H), 7.00 (d, J = 9.1 Hz, 2H), 4.14 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.47 (br, s, 2H), 3.27 (m, 5H), 2.61 (br, s, 3H), 2.48-1.85 (m, 6H) ppm.

10

Example 60

6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



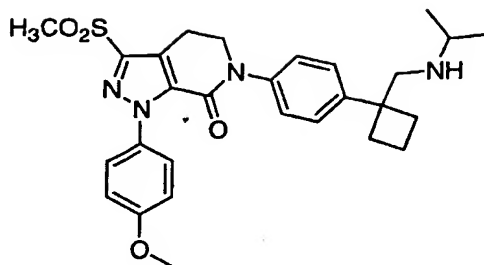
15

Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 509.4 (M+H)⁺, t_R = 4.15 min.

20

Example 61

6-(4-{1-[(isopropylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



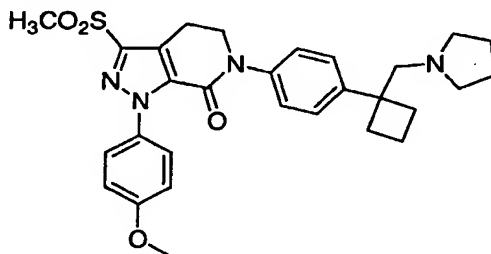
Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS

(ESI⁺) 523.4 (M+H)⁺, *t_R* = 4.27 min. ¹H NMR (acetone-*d*₆) δ 7.53 (d, *J* = 9.1 Hz, 4H), 7.36 (AA'BB', *J* = 8.4 Hz, 4H), 7.00 (d, *J* = 8.8 Hz, 2H), 4.16 (t, *J* = 6.6 Hz, 2H), 3.84 (s, 3H), 3.52 (br, s, 2H), 3.27 (m, 5H), 2.81 (m, 1H), 2.42 (m, 4H), 2.04-1.94 (m, 2H), 1.17 (d, *J* = 7.3 Hz, 6H) ppm.

10

Example 62

1-(4-methoxyphenyl)-3-(methanesulfonyl)-6-(4-[1-(1-pyrrolidinylmethyl)cyclobutyl]phenyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



15

Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (15-70% CH₃CN/H₂O in a 10-min run). LC/MS

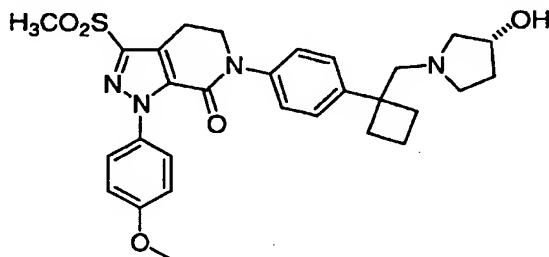
(ESI⁺) 535.4 (M+H)⁺, *t_R* = 4.74 min. ¹H NMR (acetone-*d*₆) δ

7.53 (d, *J* = 8.8 Hz, 2H), 7.41 (AA'BB', *J* = 8.8 Hz, 2H), 6.99 (d, *J* = 8.4 Hz, 2H), 4.18 (t, *J* = 6.6 Hz, 2H), 4.14 (m, 4H), 3.83 (m, 5H), 3.27 (m, 5H), 2.46 (m, 4H), 2.09-1.85 (m, 6H) ppm.

20

Example 63

6-[4-(1-[(3*R*)-3-hydroxy-1-pyrrolidinyl]methyl)cyclobutyl]phenyl]-1-(4-methoxyphenyl)-
3-(methylsulfonyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-
c]pyridin-7-one, trifluoroacetic acid salt

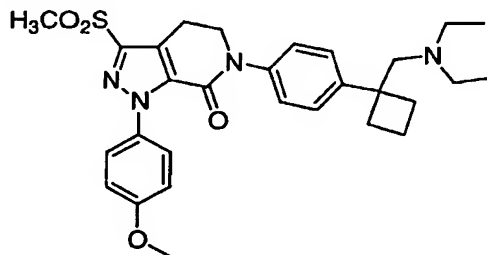


Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS

(ESI⁺) 551.4 (M+H)⁺, *t_R* = 4.06 min. ¹H NMR (acetone-*d*₆) δ 7.53 (d, *J* = 9.1 Hz, 2H), 7.40 (m, 4H), 6.99 (d, *J* = 9.1 Hz, 2H), 4.30 (m, 1H), 4.18 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.78 (m, 2H), 3.49 (m, 2H), 3.27 (m, 5H), 2.82 (m, 2H), 2.46 (m, 6H), 2.10-1.81 (m, 2H) ppm.

Example 64

6-(4-(1-[(diethylamino)methyl]cyclobutyl)phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS

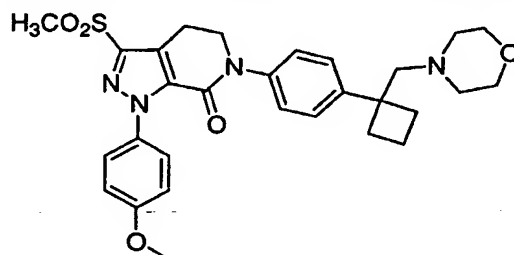
(ESI⁺) 537.4 (M+H)⁺, *t_R* = 4.62 min. ¹H NMR (acetone-*d*₆) δ 7.53 (d, *J* = 8.8 Hz, 4H), 7.41 (d, *J* = 8.4 Hz, 2H), 7.00

(d, $J = 9.2$ Hz, 2H), 4.18 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.74 (m, 2H), 3.27 (s+t, 5H), 3.03 (m, 4H), 2.49 (t, $J = 7.5$ Hz, 4H), 2.09-1.89 (m, 2H), 1.19 (t, $J = 7.4$ Hz, 6H) ppm.

5

Example 65

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



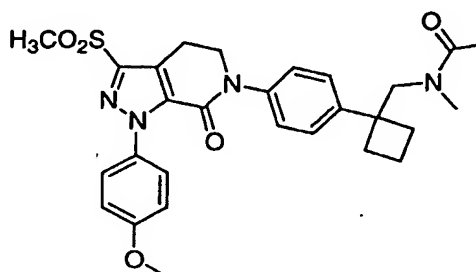
10

Following a procedure analogous to that used in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 551.4 (M+H)⁺, $t_R = 4.12$ min. ¹H NMR (acetone-*d*₆) δ 7.53 (d, $J = 8.8$ Hz, 2H), 7.48 (d, $J = 8.4$ Hz, 2H), 7.37 (d, $J = 8.8$ Hz, 2H), 7.00 (d, $J = 9.1$ Hz, 2H), 4.17 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.74 (m, 10H), 3.27 (m, 5H), 2.46 (m, 4H), 2.10-1.84 (m, 2H) ppm.

20

Example 66

N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropyl)methyl]-N-methylacetamide



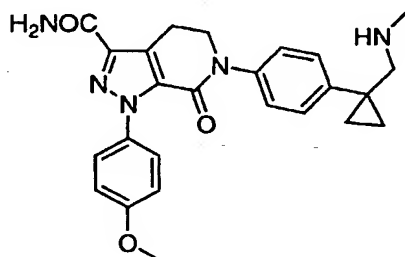
Following a procedure analogous to that of Example 21, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). ESI, LC/MS (ESI⁺) 551.4 (M+H)⁺, t_R = 4.12 min.

5

Example 67

**1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-
tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide,
trifluoroacetic acid salt**

10



Part A. 4-Iodophenylcyclopropyl acetic acid (1.93 g, 6.70 mmol) and 1-(4-methoxyphenyl)-3-(ethoxycarbonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.41 g, 4.46 mmol) were stirred in DMSO (4 mL) under N₂. K₂CO₃ (1.84 g, 13.33 mmol, 3.0 eq) was added followed by the addition of 1,10-phenanthroline (0.15 g, 20 mol%) and CuI (0.16 g, 20mol%). The resulting mixture was stirred at 110°C overnight. LC-MS showed completion of the reaction. EtOAc was added to the cooled solution. It was acidified with 1N HCl, and the organic layer was washed with H₂O, and brine, dried over MgSO₄, filtered, and concentrated to afford 1-{4-[3-(ethoxycarbonyl)-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropanecarboxylic acid (1.54 g, yield: 72.6%). LC/MS (ESI⁺) 476.4 (M+H)⁺, t_R = 2.58 min (10-90% CH₃CN/H₂O in a 4-min run).

Part B. The product from part B (1.43 g, 3.01 mmol) was stirred in THF (13 mL) at 0°C under N₂. Et₃N (0.63 mL, 4.32

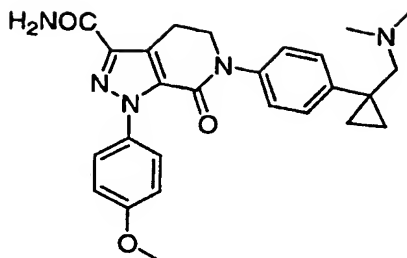
mmol, 1.5 eq) was added followed by dropwise addition of ClCOOEt (0.37 mL, 4.16 mmol, 1.3 eq). The reaction mixture was then stirred at 0°C for 20 min. TLC showed completion of the reaction. The mixture was filtered, and rinsed with anhydrous THF. The THF filtrate (ca. 20 mL) was stirred at 0°C under N₂. MeOH (3 mL) was added followed by the addition of NaBH₄ (1.03 g, 27.10 mmol, 10 eq). The resulting mixture was stirred at 0°C for 15 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give 6-{4-[1-(hydroxymethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-(ethoxycarbonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.30 g, 99%). LC/MS (ESI⁺) 462.6 (M+H)⁺, t_R = 2.57 min (10-90% CH₃CN/H₂O in a 4-min run).

Part C. The product from part B (1.90 g, 4.12 mmol) was stirred in anhydrous CH₂Cl₂ (13 mL) at RT under N₂. NaOAc (1.01 g, 12.20 mmol, 3 eq) and molecular sieves (2.0 g) were added followed by the addition of PCC (1.78 g, 8.24 mmol, 2 eq). The resulting slurry was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was filtered, and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to afford ethyl 6-[4-(1-formylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (1.90 g, yield: 100%). LC/MS (ESI⁺) 460.6 (M+H)⁺, t_R = 2.69 min (10-90% CH₃CN/H₂O in a 4-min run).

- Part D. The product from part C (250 mg, 0.55 mmol), methylamine hydrochloride (0.5 g, excess) were stirred in dichloroethane (15 mL) at RT under N₂. NaBH(OAc)₃ (1.03 g, 4.86 mmol) was added followed by addition of three drops of HOAc. The reaction mixture was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was quenched with H₂O, and extracted with EtOAc (2x). The organic layer was washed with brine, dried over MgSO₄, filtered, and concentrated to dryness to obtain crude ethyl 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (109 mg, yield: 42%). LC/MS (ESI⁺) 475.4 (M+H)⁺, t_R = 2.08 min.
- Part E. The product from part D (50 mg, 0.105 mmol) was stirred in ethylene glycol (saturated with NH₃) in a capped Pyrex tube at 80°C for 4 h. After cooling, the mixture was diluted with MeOH, and purified by prep LC-MS (5-98% CH₃CN in H₂O in a 10-min run) to afford 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide (29 mg, yield: 60%). LC/MS (ESI⁺) 445.4 (M+H)⁺, t_R = 3.53 min. ¹H NMR (acetone-d₆) δ 7.49 (m, 4H), 7.29 (d, J = 8.1 Hz, 2H), 6.97 (d, J = 9.1 Hz, 2H), 4.06 (t, J = 6.6 Hz, 2H), 3.82 (s, 3H), 3.32 (s, 2H), 3.24 (t, J = 6.6 Hz, 2H), 2.66 (s, 3H), 1.10 (m, 2H), 0.94 (m, 2H) ppm.

Example 68

- 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt



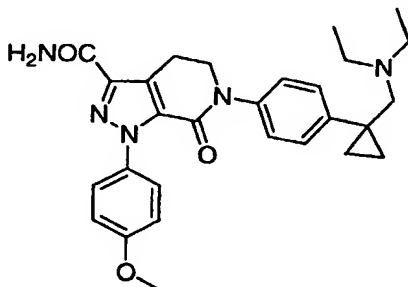
Following a procedure analogous to that used in Example 67, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS

(ESI⁺) 460.6 (M+H)⁺, *t_R* = 3.93 min. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.33 (m, 2H), 6.97 (d, *J* = 9.2 Hz, 2H), 4.10 (t, *J* = 6.3 Hz, 2H), 3.82 (s, 3H), 3.52 (m, 2H), 3.26 (t, *J* = 6.3 Hz, 2H), 2.69 (m, 6H), 1.18 (m, 2H), 1.04 (m, 2H) ppm.

10

Example 69

6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt



15

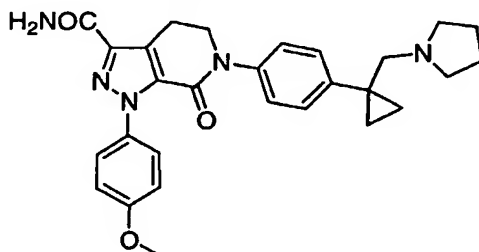
Following a procedure analogous to that used in Example 67, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS

(ESI⁺) 488.6 (M+H)⁺, *t_R* = 3.90 min. ¹H NMR (acetone-*d*₆) δ 7.57 (d, *J* = 8.8 Hz, 2H), 7.51 (d, *J* = 8.8 Hz, 2H), 7.34 (d, *J* = 8.8 Hz, 2H), 6.97 (d, *J* = 9.1 Hz, 2H), 4.08 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.62 (m, 2H), 3.24 (m, 6H), 1.16 (m, 8H), 1.05 (m, 2H) ppm.

20

Example 70

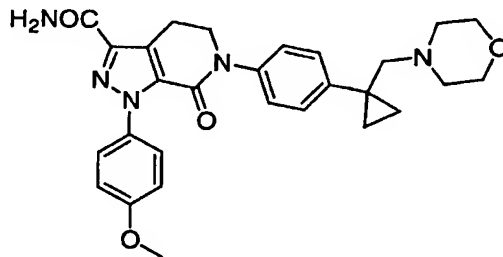
1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 67, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 486.4 (M+H)⁺, t_R = 3.88 min. ¹H NMR (acetone-d₆) δ 7.52 (m, 2H), 7.32 (m, 4H), 6.99 (m, 2H), 4.09 (m, 2H), 3.82 (s, 3H), 3.56 (m, 6H), 3.26 (m, 2H), 1.91 (m, 4H), 1.13 (m, 2H), 0.98 (m, 2H) ppm.

Example 71

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt



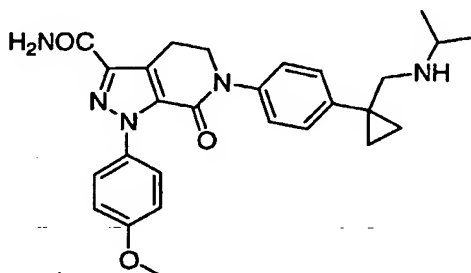
Following a procedure analogous to that used in Example 67, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 3.67 min). LC/MS (ESI⁺) 502.6 (M+H)⁺. ¹H NMR (acetone-d₆) δ 7.51 (d, J = 8.6 Hz, 4H), 7.34 (d, J = 8.4 Hz, 2H), 7.25 (d, J = 8.4

Hz, 4H), 6.95 (d, $J = 9.2$ Hz, 2H), 4.09 (t, $J = 6.6$ Hz, 2H), 3.82 (s, 3H), 3.48 (m, 6H), 3.26 (t, $J = 6.6$ Hz, 2H), 2.82 (m, 2H), 2.40 (m, 2H), 0.81 (m, 2H), 0.73 (m, 2H) ppm.

5

Example 72

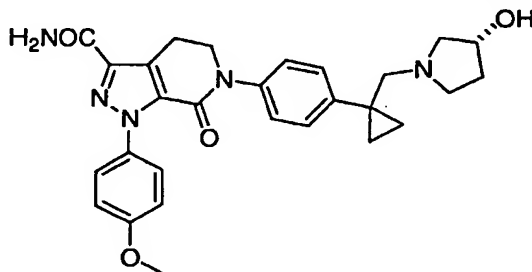
6-(4-(1-[(isopropylamino)methyl]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt



10 Following a procedure analogous to that used in Example 67, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, $t_R = 4.18$ min). LC/MS (ESI⁺) 474.4 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.53 (m, 4H), 7.28 (d, $J = 8.1$ Hz, 2H), 6.97 (d, $J = 7.7$ Hz, 2H),
 15 4.08 (m, 2H), 3.82 (s, 3H), 3.35 (m, 3H), 3.25 (m, 2H), 1.27 (d, $J = 6.2$ Hz, 6H), 1.11 (m, 2H), 0.92 (m, 2H) ppm.

Example 73

20 **6-[4-(1-[(3R)-3-hydroxy-1-pyrrolidinyl]methyl]cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide, trifluoroacetic acid salt**



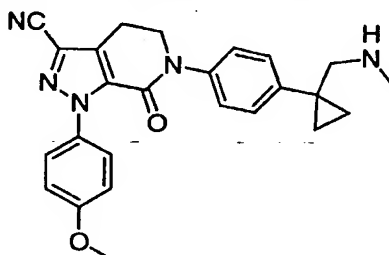
Following a procedure analogous to that used in Example 67, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 502.4 (M+H)⁺, t_R = 3.81 min.

5

Example 74

1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-
tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile,
trifluoroacetic acid salt

10



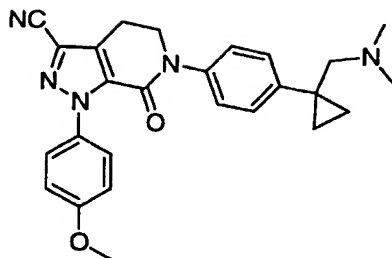
The product of Example 67 (15 mg) was stirred in DMF (0.5 mL) at RT in a capped vial. Two drops of thionyl chloride was added. The reaction was completed in 10 min. The mixture was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run) to give pure 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile (11 mg, yield: 76.2%). LC/MS (ESI⁺) 428.4 (M+H)⁺, t_R = 4.49 min.

20

Example 75

6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt

25



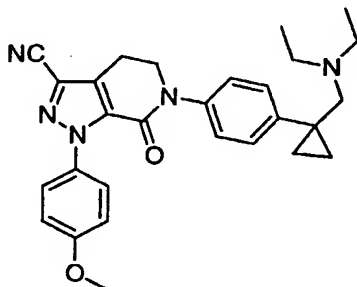
Following a procedure analogous to that used in Example 74, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 4.44 min).

5 LC/MS (ESI⁺) 442.6 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.53 (m, 4H), 7.33 (d, J = 8.4 Hz, 2H), 6.99 (d, J = 9.2 Hz, 2H), 4.18 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.57 (s, 2H), 3.18 (t, J = 6.6 Hz, 2H), 2.78 (s, 6H), 1.16 (m, 2H), 1.06 (m, 2H) ppm.

10

Example 76

6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



15

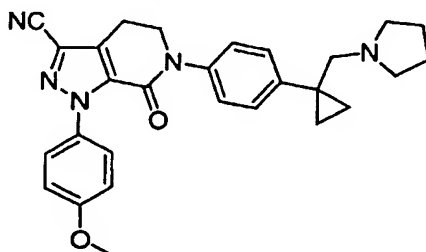
Following a procedure analogous to that used in Example 74, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 4.60 min).

LC/MS (ESI⁺) 470.6 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.58 (d, J = 8.4 Hz, 2H), 7.54 (d, J = 8.4 Hz, 2H), 7.35 (d, J = 8.4 Hz, 2H), 6.99 (d, J = 9.1 Hz, 2H), 4.19 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.58-3.18 (m, 6H), 3.19 (t, J = 6.6 Hz, 2H), 1.18 (m, 8H), 1.06 (m, 2H) ppm.

20

Example 77

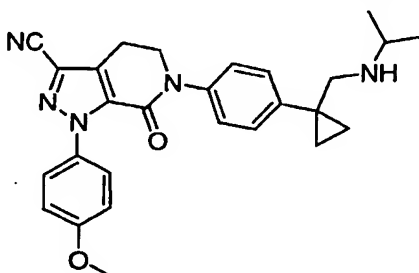
1-(4-methoxyphenyl)-7-oxo-6-(4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



Following a procedure analogous to that used in step F of Example 74, the title compound was prepared. LC/MS (ESI⁺) 468.4 (M+H)⁺, t_R = 4.49 min. ¹H NMR (acetone-*d*₆) δ 7.52 (d, J = 9.0 Hz, 2H), 7.44 (AA'BB', J = 8.6 Hz, 4H), 6.99 (d, J = 8.8 Hz, 2H), 4.18 (t, J = 6.6 Hz, 2H), 3.83 (s, 3H), 3.59 (m, 2H), 3.19 (t, J = 6.6 Hz, 2H), 2.75 (m, 4H), 2.01 (m, 4H), 1.14 (m, 2H), 1.00 (m, 2H).

Example 78

6-(4-(1-[(isopropylamino)methyl]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



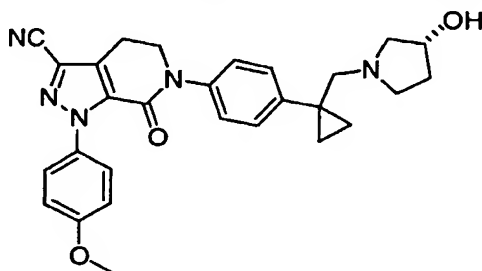
Following a procedure analogous to that used in Example 74, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 4.57 min). LC/MS (ESI⁺) 456.6 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.53 (m, 4H), 7.31 (d, J = 8.4 Hz, 2H), 7.00 (d, J = 9.1 Hz, 2H),

4.17 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.19 (t, $J = 6.6$ Hz, 2H), 3.17 (m, 3H), 1.28 (d, $J = 6.6$ Hz, 6H), 1.13 (m, 2H), 0.93 (m, 2H) ppm.

5

Example 79

6-[4-(1-[(3*R*)-3-hydroxy-1-pyrrolidinyl)methyl]cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile, trifluoroacetic acid salt



10

Following a procedure analogous to that used in Example 74, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, $t_R = 4.34$ min).

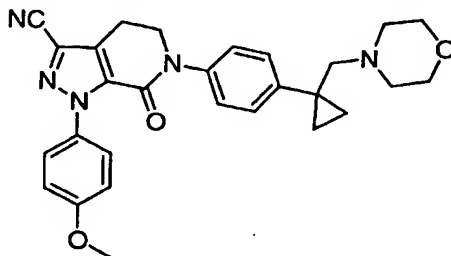
LC/MS (ESI⁺) 484.4 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.53 (m,

15 4H), 7.35 (d, $J = 8.4$ Hz, 2H), 6.99 (d, $J = 9.2$ Hz, 2H), 4.19 (t, $J = 6.6$ Hz, 2H), 3.83 (s, 3H), 3.62 (m, 5H), 3.19 (t, $J = 6.6$ Hz, 2H), 2.91 (m, 2H), 1.85 (m, 2H), 1.18 (m, 2H), 1.01 (m, 2H) ppm.

20

Example 80

1-(4-methoxyphenyl)-6-[4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl]-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile, trifluoroacetic acid salt



25

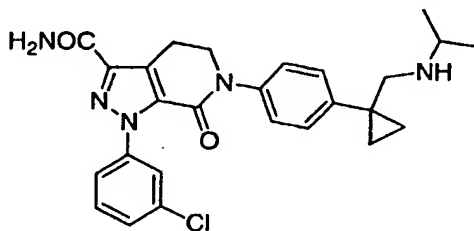
Following a procedure analogous to that used in Example 67, the title compound was prepared. It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, *t_R* = 4.46 min).

LC/MS (ESI⁺) 484.6 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.52 (m, 4H), 7.32 (d, *J* = 8.4 Hz, 2H), 6.99 (d, *J* = 9.2 Hz, 2H), 4.17 (t, *J* = 6.6 Hz, 2H), 3.83 (s, 3H), 3.47 (m, 7H), 3.62 (s, 2H), 3.49 (m, 2H), 3.17 (t, *J* = 6.6 Hz, 2H), 3.07 (m, 2H), 1.17 (m, 2H), 1.06 (m, 2H) ppm.

10

Example 81

1-(3-chlorophenyl)-6-{4-[1-(isopropylamino)methyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



15 Part A. 1-(3-Chlorophenyl)-3-(ethoxycarbonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (1.14 g, 3.57 mmol) and 4-iodophenylcyclopropyl acetic acid (1.13 g, 1.1 eq) were stirred in DMSO (4 mL) under N₂. K₂CO₃ (1.48 g, mmol, 3 eq) was added, followed by the addition of 1,10-phenanthroline (0.13 g, 20 mol%) and CuI (0.14 g, 20 mol%).
20 The resulting mixture was stirred at 130°C overnight. LC-MS showed completion of the reaction. EtOAc was added to the cooled solution. It was washed with 1N HCl, H₂O, and brine; dried over MgSO₄; filtered; and concentrated in
25 vacuo to give almost pure 1-{4-[1-(3-chlorophenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropanecarboxylic acid (0.87 g, yield: 51%). LC/MS (ESI⁺) 480.4 (M+H)⁺.

Part B. The product from Part A (0.54 g, 1.13 mmol) was stirred in THF (6 mL) at 0°C under N₂. Et₃N (0.24 mL, 1.5 eq) was added, followed by dropwise addition of ClCOOEt (0.14 mL, 1.3 eq). The reaction mixture was then stirred
5 at 0°C for 1 h. TLC showed the completion of the reaction. The mixture was filtered through a filter funnel and rinsed with anhydrous THF. The THF filtrate (ca. 10 mL) was stirred at 0°C under N₂. NaBH₄ (0.52 g, 10 eq) was added, followed by the addition of MeOH (2.5 mL). The resulting
10 mixture was stirred at 0°C. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give 1-(3-
15 chlorophenyl)-6-{4-[1-(hydroxymethyl)cyclopropyl]phenyl}-3-(ethoxycarbonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.31 g, yield: 52.4%). LC/MS (ESI⁺) 466.4 (M+H)⁺.

20 Part C. The product from Part B (0.31 g, 0.22 mmol) was stirred in anhydrous CH₂Cl₂ (5 mL) at RT under N₂. NaOAc (0.16 g, 1.95 mmol) and molecular sieves (0.5 g) were added, followed by the addition of PCC (0.29 g, 1.34 mmol). The resulting slurry was stirred at RT for 1.5h.
25 Analytical LC-MS showed completion of the reaction. The mixture was filtered through Celite, and rinsed with CH₂Cl₂. The filtrate was washed with H₂O (2x), brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness to give almost pure 1-(3-chlorophenyl)-6-[4-(1-
30 (formylcyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.20 g, yield: 87.4%). LC/MS (ESI⁺) 464.4 (M+H)⁺.

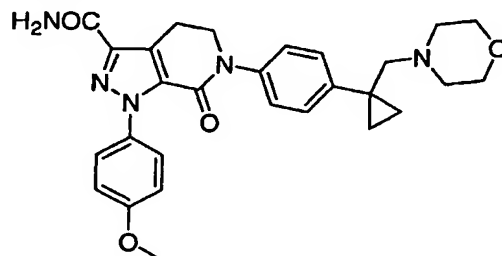
Part D. The product from Part C (100 mg) and isopropyl
35 amine (0.1 mL, excess) were stirred in dichloroethane (1

mL) in a capped vial. $\text{NaBH}(\text{OAc})_3$ (200 mg) was added, followed by addition of one drop of HOAc. The reaction mixture was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was
 5 evaporated, and dissolved in aqueous MeOH. It was then purified by prep LC-MS (5-98% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 10-min run) to obtain pure 1-(3-chlorophenyl)-6-{4-[1-(isopropylamino)methyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide (60 mg,
 10 yield: 55%). LC/MS (ESI^+) 507.4 ($\text{M}+\text{H}^+$), $t_R = 4.68$ min.

Part E. The product from part D (60 mg) was stirred in ethylene glycol (saturated with NH_3) in a capped Pyrex tube at 80°C for 4 h. After cooling, the mixture was diluted
 15 with MeOH and purified by prep LC-MS (5-98% CH_3CN in H_2O in a 10-min run) to afford the title compound (35 mg, yield: 62%). LC/MS (ESI^+) 478.4 ($\text{M}+\text{H}^+$), $t_R = 4.34$ min. ^1H NMR (acetone- d_6) δ 7.74 (s, 1H), 7.63 (m, 1H), 7.45 (m, 4H), 7.30 (d, $J = 8.4$ Hz, 2H), 4.11 (t, $J = 6.6$ Hz, 2H), 3.44
 20 (m, 1H), 3.38 (m, 2H), 3.26 (t, $J = 6.6$ Hz, 2H), 1.29 (d, $J = 6.6$ Hz, 6H), 1.12 (m, 2H), 0.94 (m, 2H) ppm.

Example 82

1-(3-chlorophenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide,
 25 trifluoroacetic acid salt

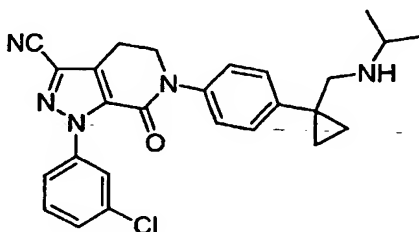


Following a procedure analogous to that used in Example 81,
 30 the title compound was prepared. The product was purified

by prep LC-MS (5-98% CH₃CN in H₂O in a 10-min run). LC/MS
 (ESI⁺) 506.6 (M+H)⁺, t_R = 4.57 min. ¹H NMR (acetone-d₆) δ
 7.73 (s, 1H), 7.63 (m, 1H), 7.47 (m, 4H), 7.29 (m, 2H),
 4.09 (t, J = 6.6 Hz, 2H), 3.74 (m, 4H), 3.53 (m, 2H), 3.27
 5 (m, 2H), 3.07 (m, 4H), 1.07 (m, 2H), 1.00 (m, 2H) ppm.

Example 83

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-
 methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
 10 c]pyridine-3-carbonitrile, trifluoroacetic acid salt

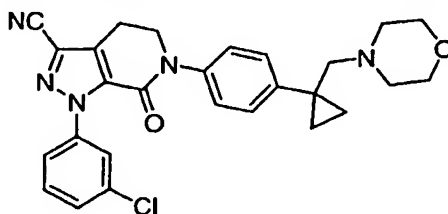


Following a procedure analogous to that used in step F of
 Example 74, the title compound was prepared. It was then
 purified by prep LC-MS (35-98% CH₃CN/H₂O in a 10-min run, t_R
 15 = 2.78 min). LC/MS (ESI⁺) 460.6 (M+H)⁺. ¹H NMR (acetone-
 d₆) δ 7.75 (s, 1H), 7.63 (m, 1H), 7.53 (m, 4H), 7.31 (d, J =
 8.8 Hz, 2H), 4.19 (t, J = 6.6 Hz, 2H), 3.48 (m, 1H), 3.41
 (m, 2H), 3.21 (t, J = 6.6 Hz, 2H), 1.29 (d, , J = 6.3 Hz,
 6H), 1.14 (m, 2H), 0.95 (m, 2H) ppm.

20

Example 84

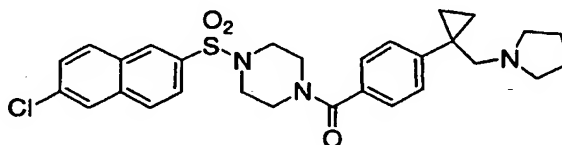
1-(4-methoxyphenyl)-6-{4-[1-(4-
 morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-
 tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile,
 25 trifluoroacetic acid salt



Following a procedure analogous to that used in Example 74, the title compound was prepared. It was then purified by prep LC-MS (35-98% CH₃CN/H₂O in a 10-min run, t_R = 2.80 min). LC/MS (ESI⁺) 488.6 (M+H)⁺. ¹H NMR (acetone-*d*₆) δ 7.73 (s, 1H), 7.62 (m, 1H), 7.54 (m, 4H), 7.34 (m, 2H), 4.19 (m, 2H), 3.83 (m, 8H), 3.61 (m, 2H), 3.19 (m, 2H), 1.17 (m, 2H), 1.05 (m, 2H) ppm.

Example 85

1-[(6-chloro-2-naphthyl)sulfonyl]-4-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl}piperazine



Part A. 1-(4-Iodophenyl)cyclopropane carboxylic acid (0.64 g, 2.25 mmol) was stirred in THF (10 mL) at 0°C under N₂. Et₃N (0.47 mL, 3.37 mmol) was added, followed by dropwise addition of ClCO₂Et (0.28 mL, 2.93 mmol). The reaction mixture was then stirred at 0°C for 30 min. TLC showed the completion of the reaction. The mixture was filtered through a filter funnel and rinsed with anhydrous THF. The THF filtrate (ca.15 mL) was stirred at 0°C under N₂. NaBH₄ (0.41 g, 10.8 mmol) was added, followed by addition of MeOH (3 mL). The resulting mixture was stirred at 0°C for 30 min. Analytical LC-MS showed completion of the reaction. Sat'd Na₂SO₄ was then added. The mixture was extracted with EtOAc (2x). The organic layer was washed with H₂O (2x) and brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness. The resulting alcohol was stirred in anhydrous CH₂Cl₂ (10 mL) at RT under N₂. NaOAc (0.42 g, 5.12 mmol) and molecular sieves (4Å, 0.75 g) were added, followed by the addition of PCC (0.83 g, 3.84 mmol). The resulting slurry was stirred at RT for 1.5 h. Analytical LC-MS showed completion of the reaction. The mixture was

filtered through Celite, and rinsed with CH_2Cl_2 . The filtrate was washed with H_2O (2x) and brine (2x), dried over Na_2SO_4 , filtered, and concentrated in vacuo to give almost pure 4-iodophenylcyclopropanecarbaldehyde. This aldehyde and pyrrolidine (0.37 mmol) were stirred in dichloroethane (6 mL) at RT under N_2 . $\text{NaBH}(\text{OAc})_3$ (1.37 mg, mmol) was added, followed by addition of several drops of HOAc. The reaction mixture was stirred at RT for 20 min. Analytical LC-MS showed completion of the reaction. H_2O was added. The mixture was extracted with EtOAc; and the organic extracts were washed with H_2O (2x) and brine (2x), dried over Na_2SO_4 , filtered, and concentrated to dryness to give almost pure 1-([1-(4-iodophenyl)cyclopropyl]methyl)pyrrolidine (0.41 g, yield % for 3 steps). LC/MS (ESI^+) 328.2 ($\text{M}+\text{H}^+$) (10-90% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 4-min run, $t_R = 1.77$ min).

Part B. The product from part A (0.40 g, 1.24 mmol), KOAc (0.61 g, 5.0 eq), $\text{Pd}(\text{OAc})_2$ (0.03 g, 0.1 eq), and dppf (0.14 g, 0.2 eq) were stirred in DMF (3 mL) at RT. The mixture was degassed twice and purged with CO. The mixture was heated at 60°C under CO atmosphere with a balloon for 2.5 h. LS-MS showed completion of the reaction. After cooling, H_2O was added. The mixture was extracted with EtOAc (2x). The aqueous layer was then acidified, and concentrated to dryness. MeOH was added, and filtered off inorganic salts. The filtrate was concentrated and vacuum dried to give almost pure 4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoic acid (0.32 g, yield: 96%). LC/MS (ESI^+) 246.4 ($\text{M}+\text{H}^+$) (10-90% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 4-min run, $t_R = 1.32$ min).

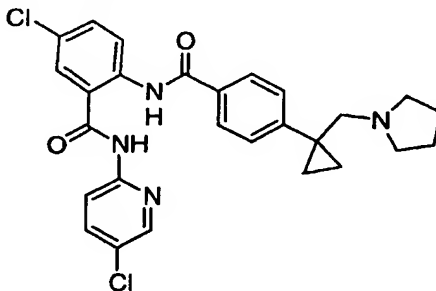
Part C. The product of part B (0.41 g, 1.68 mmol) was stirred in CH_2Cl_2 (10 mL) at RT under N_2 . $(\text{COCl})_2$ (0.5 mL)

was added, followed by the addition of one drop of DMF. The mixture was stirred at RT for 1 h. The solvent was evaporated and dried in vacuo. The resulting acid chloride (0.16 g, 0.61 mmol) was dissolved in CH₂Cl₂ (10 mL), 1-[(6-chloro-2-naphthyl)sulfonyl]piperazine (0.21 g, 0.61 mmol) was added, followed by the addition of DIEA (0.21 mL, 1.21 mmol). The resulting mixture was stirred at RT for 20 min. Analytical LC-MS showed completion of the reaction. The solvent was evaporated. The residue was dissolve in MeOH, and purified by RP Prep LC-MS (5-98% CH₃CN in H₂O in a 10-min run) to give pure title compound (210 mg, yield: 64.1%). It was then purified by prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run, t_R = 4.71 min). LC/MS (ESI⁺) 538.4 (M+H)⁺.

15

Example 86

5-chloro-N-(5-chloro-2-pyridinyl)-2-((4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl)amino)benzamide, trifluoroacetic acid salt



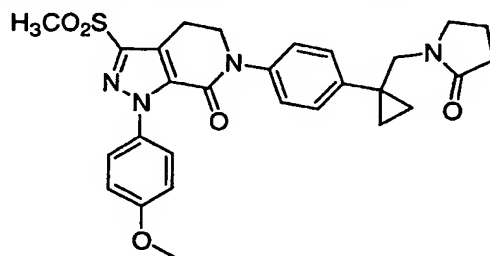
Part A. The product from Part B of Example 85 (0.16 g, 0.65 mmol) was stirred in CH₂Cl₂ (5 mL) at RT under N₂. (COCl)₂ (0.2 mL) was added. The mixture was stirred at RT for 1 h. The solvent was evaporated and dried in vacuo. The resulting acid chloride was dissolved in CH₂Cl₂ (6 mL), 2-amino-5-chlorobenzoic acid methyl ester (0.16 g, 0.86 mmol) was added, followed by the addition of DIEA (0.30 mL). The resulting mixture was stirred at RT for 2 h. Analytical LC-MS showed completion of the reaction. The solvent was evaporated. The residue was dissolve in EtOAc,

washed with H₂O, brine, dried over MgSO₄, and concentrated to give methyl 5-chloro-2-({4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl}amino)benzoate (55 mg, yield: 21%). LC/MS (ESI⁺) 413.4 (M+H)⁺, t_R = 2.19 min (10-90% CH₃CN/H₂O in a 4-min run).

Part B. The product from Part A (30 mg) and 5-chloro-2-aminopyridine (14 mg) were stirred in CH₂Cl₂ (1 mL) at RT under N₂. Me₃Al in toluene (0.45 mL, 0.23 mmol) was added dropwise. The resulting solution was stirred at RT for 1h and at reflux for 2h. The solvent was evaporated after cooling. The residue was dissolved in MeOH, and purified by LC-MS (5-98% CH₃CN in H₂O in a 10-min run) to give pure 5-chloro-N-(5-chloro-2-pyridinyl)-2-({4-[1-(1-pyrrolidinylmethyl)cyclopropyl]benzoyl}amino)benzamide (6 mg, yield: 16%). LC/MS (ESI⁺) 509.2 (M+H)⁺, t_R = 2.21 min (10-90% CH₃CN/H₂O in a 4-min run).

Example 87

20 1-(4-Methoxyphenyl)-3-methanesulfonyl-6-({4-[1-(2-oxo-pyrrolidin-1-ylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Part A. 1-(1-Bromomethylcyclopropyl)-4-iodobenzene (2.0 g, 5.97 mmol) and NaN₃ (1.0 g, 15.38 mmol, 2.6 eq) were stirred in DMF (10 mL) overnight. Analytical LC-MS showed completion of the reaction. EtOAc was added to the solution. The mixture was washed with H₂O and brine, dried over MgSO₄, and concentrated to give 1-(1-azidomethylcyclopropyl)-4-iodobenzene (1.43 g, yield: 80%). The azide

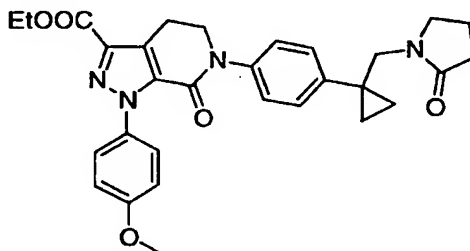
- (1.40 g, 4.68 mmol) and PPh_3 (1.84 g, 7.02 mmol, 1.5 eq) were stirred in THF (10 mL) at RT for 40 min. H_2O (2 mL) was added, and the solution was stirred at 50°C for 6h. LC-MS showed completion of the reaction. The mixture was
5 extracted with Et_2O (2x). The aqueous layer was basified with 50% NaOH, extracted with CH_2Cl_2 (2x), washed with H_2O , brine, dried over MgSO_4 , and concentrated to give 1-(4-iodophenyl)cyclopropyl methylamine (0.98 g, yield: 75%).
- 10 Part B. The product from Part A (0.36 g, 1.31 mmol) was stirred in dry CH_2Cl_2 (10 mL) at RT. NaOH (0.16 g, 3.93 mmol, 3 eq) was added, followed by the addition of 4-chlorobutyryl chloride (0.16 mL, 1.42 mmol). The reaction mixture was stirred at RT for 1h. It was washed with H_2O
15 and brine, dried over MgSO_4 , and concentrated to dryness. The residue was dissolved in THF (10 mL). K-O-tBu (0.29 g, 2.62 mmol) was added as one single portion. The mixture was stirred at 0°C under N_2 for 1h. LC-MS showed completion of the reaction. EtOAc was added. It was washed with H_2O
20 and brine, dried over MgSO_4 , and concentrated to produce 1-[1-(4-iodophenyl)cyclopropylmethyl]-pyrrolidin-2-one (0.36 g, yield: 86%). LC/MS (ESI^+) 342.0 ($\text{M}+\text{H}^+$), $t_R = 2.86$ min (10-90% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ in a 4-min run).
- 25 Part C. The product from Part B (0.18 g, 0.56 mmol) and 1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.16 g, 0.47 mmol) were stirred in DMSO (1 mL) under N_2 . K_2CO_3 (0.20 g, 1.44 mmol) was added, followed by the addition of CuI (0.030 g, 20
30 mol%) and 1,10-phenanthroline (0.028 g, 20 mol%). The resulting mixture was heated at 120°C overnight. After cooling, it was extracted with EtOAc (2x), washed with H_2O and brine, dried over MgSO_4 , filtered, and concentrated to dryness. The residue was purified by flash column

chromatography (silica gel, CH₂Cl₂:EtOAc = 1:1, then EtOAc) to give the desired compound (83 mg, yield: 25%). LC/MS (ESI⁺) 535.2 (M+H)⁺, t_R = 3.45 min (10-90% CH₃CN/H₂O in a 6-min run). ¹H NMR (CDCl₃) δ 7.45 (d, J = 8.8 Hz, 2H), 7.25 (AA'BB', J = 8.6 Hz, 4H), 6.91 (d, J = 9.2 Hz, 2H), 4.10 (t, J = 6.6 Hz, 2H), 3.80 (s, 3H), 3.68 (s, 2H), 3.46 (m, 2H), 3.29 (m, 3H), 3.23 (t, J = 6.6 Hz, 2H), 2.25 (m, 2H), 1.86 (m, 2H), 0.86 (m, 4H) ppm.

10

Example 88

1-(4-Methoxyphenyl)-7-oxo-6-(4-[1-(2-oxo-pyrrolidin-1-ylmethyl)cyclopropyl]phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester

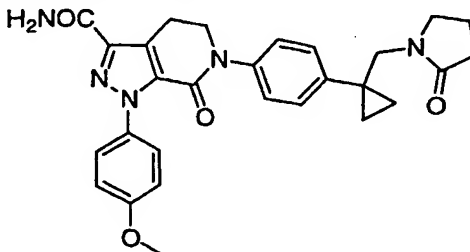


15 Following a procedure analogous to that used for the preparation of Example 87, the title compound was prepared. The product was purified by silica gel column chromatography. LC/MS (ESI⁺) 529.4 (M+H)⁺, t_R = 3.14 min (25-90% CH₃CN/H₂O in a 6-min run).

20

Example 89

1-(4-Methoxyphenyl)-7-oxo-6-(4-[1-(2-oxo-pyrrolidin-1-ylmethyl)cyclopropyl]phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide

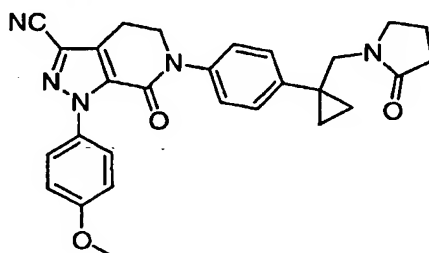


25

Following a procedure analogous to that used for the preparation of Example 67, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 500.2 (M+H)⁺, t_R = 3.28 min
5 (10-90% CH₃CN/H₂O in a 6-min run).

Example 90

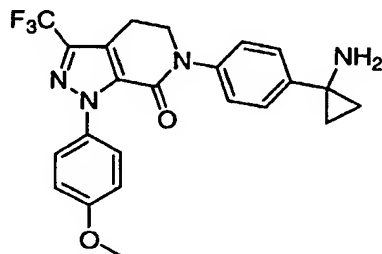
1-(4-Methoxyphenyl)-7-oxo-6-(4-[1-(2-oxo-pyrrolidin-1-ylmethyl)cyclopropyl]phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile
10



Following a procedure analogous to that used for the preparation of Example 74, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). Analytical LC/MS (ESI⁺) 482.4 (M+H)⁺, t_R = 2.63 min (35-95% CH₃CN/H₂O in a 6-min run).
15

Example 91

6-[4-(1-Aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridine-7-one, trifluoroacetic acid salt
20

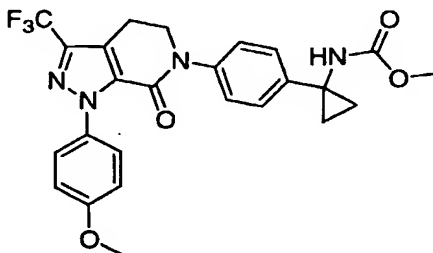


The product of Part D in Example 1 (ca. 0.50 g) was stirred in dry toluene at RT. DPPA (0.25 mL) was added, followed by the addition of Et₃N (0.35 mL). The resulting mixture
25

was stirred at 100°C for 3h. After cooling to RT, 8N HCl (10 mL) was added. The resulting mixture was heated at 100°C overnight. The cooled mixture was extracted with Et₂O (2x). The aqueous layer was basified with 50% NaOH. The mixture was extracted with chloroform (2x). The organics were washed with H₂O and brine, dried over MgSO₄, and concentrated to dryness. The residue was dissolved in MeOH, and purified by prep LC/MC (5-98% CH₃CN/H₂O in a 10-min run) to give the desired product. Analytical LC/MS (ESI⁺) 443.2 (M+H)⁺, t_R = 2.69 min (35-95% CH₃CN/H₂O in a 6-min run).

Example 92

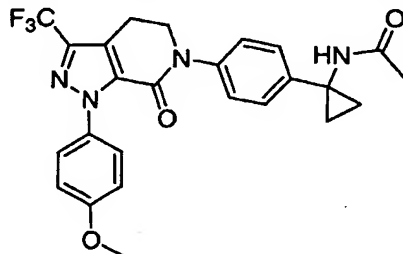
(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-carbamic acid methyl ester



Following a procedure analogous to that used for the preparation of Example 91, the title compound was prepared by using MeOH instead of conc. HCl as the solvent. Silica gel purification yielded the pure desired product. LC/MS (ESI⁺) 501.6 (M+H)⁺, t_R = 3.19 min (35-95% CH₃CN/H₂O in a 6-min run). ¹H NMR (CDCl₃) δ 7.46 (d, J = 8.8 Hz, 2H), 7.24 (m, 4H), 6.91 (d, J = 9.2 Hz, 2H), 4.12 (t, J = 6.6 Hz, 2H), 3.81 (s, 3H), 3.64 (s, 2H), 3.14 (t, J = 6.6 Hz, 2H), 1.22 (m, 4H) ppm.

Example 93

***N*-(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-acetamide**



5

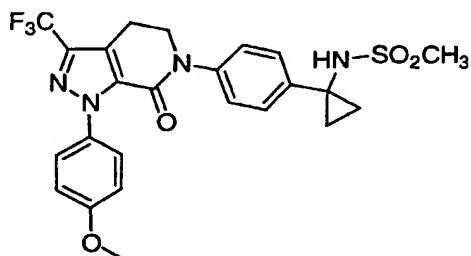
Following a procedure analogous to that used for the preparation of Example 21, the title compound was prepared. Silica gel purification yielded the pure desired product.

LC/MS (ESI⁺) 485.2 (M+H)⁺, *t_R* = 3.06 min (35-95% CH₃CN/H₂O in a 6-min run). ¹H NMR (CDCl₃) δ 7.46 (d, *J* = 8.8 Hz, 2H), 7.24 (d, *J* = 8.8 Hz, 2H), 6.94 (AA'BB', *J* = 9.1 Hz, 4H), 4.09 (m, 2H), 3.81 (m, 2H), 3.14 (t, *J* = 6.6 Hz, 2H), 2.38 (s, 3H), 1.57 (m, 2H), 1.40 (m, 2H) ppm.

15

Example 94

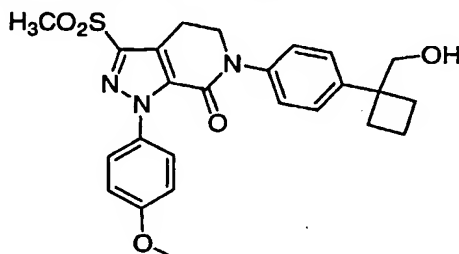
***N*-(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-methanesulfonamide**



20 Silica gel purification yielded the pure desired product. LC/MS (ESI⁺) 521.2 (M+H)⁺, *t_R* = 3.14 min (35-95% CH₃CN/H₂O in a 6-min run). ¹H NMR (CDCl₃) δ 7.46 (d, *J* = 9.2 Hz, 2H), 7.35 (AA'BB', *J* = 8.8 Hz, 4H), 6.92 (d, *J* = 8.8 Hz, 2H), 5.19 (s, 1H), 4.13 (t, *J* = 6.6 Hz, 2H), 3.81 (s, 3H), 3.16 (t, *J* = 6.6 Hz, 2H), 1.26 (m, 2H), 1.18 (m, 2H) ppm.

Example 95

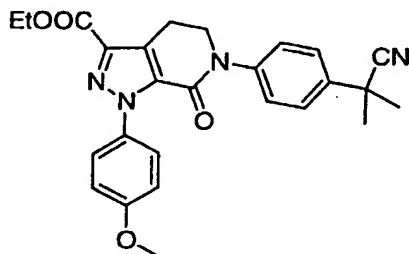
6-[4-(1-Hydroxymethylcyclopropyl)phenyl]-3-(methanesulfonyl)-1-(4-methoxyphenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Following a procedure analogous to that used for the preparation of product of Part C in Example 48, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). LC/MS (ESI⁺) 481.4 (M+H)⁺, t_R = 5.51 min. ¹H NMR (acetone-d₆) δ 7.53 (d, J = 8.8 Hz, 2H), 7.27 (d, J = 8.8 Hz, 2H), 7.14 (d, J = 8.8 Hz, 2H), 6.98 (d, J = 8.8 Hz, 2H), 4.14 (t, J = 6.6 Hz, 2H), 3.83 (m, 4H), 3.63 (s, 2H), 3.27 (t, J = 6.6 Hz, 2H), 3.26 (s, 3H), 2.02 (m, 4H), 1.81 (m, 2H) ppm.

Example 96

Ethyl 6-[4-(cyano-dimethyl-methyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate



Part A. To 4-iodobenzylbromide (25 g, 84 mmol) in boiling EtOH (100 mL) was added potassium cyanide (8.2 g, 126 mmol) through the condenser. The reaction was heated 24h, then

cooled and EtOH removed. The aqueous layer was extracted with EtOAc and dried (Na_2SO_4) to afford crude 4-iodobenzyl nitrile. The 4-iodobenzyl nitrile was first treated with HCl gas in MeOH to afford conversion to the ester. The mixture was concentrated in vacuo and treated with MeOH (4.7 mL) and chlorotrimethylsilane (10.7 mL) at 50°C for 4h. The reaction was cooled and quenched with H_2O (3.5 mL). Dichloromethane (150 mL) was added followed by Na_2CO_3 (8.9 g) and the mixture was stirred at room temperature for 1h. The organics were separated and dried (Na_2SO_4), filtered, and concentrated to afford 21 g crude (4-iodo-phenyl)-acetic acid methyl ester. ^1H NMR (CDCl_3) δ 7.66 (d, $J = 8.4$ Hz, 2H), 7.04 (d, $J = 8.4$ Hz, 2H), 3.69 (s, 3H), 3.56 (s, 2H) ppm.

15

Part B. To a THF (100 mL) solution containing sodium hydride (9.5 g, 0.23 mol) at 0°C was added dropwise crude methyl-(4-iodo-phenyl)-acetic acid methyl ester (21 g, 79 mmol, from Part A in THF (50 mL). After the addition was complete, methyl iodide (11.4 mL, 0.18 mol) in THF (20 mL) was added and the reaction was stirred 72h at rt. The reaction mixture was quenched with ice water followed by extraction with EtOAc. Drying with Na_2SO_4 afforded 27 g of a crude mixture of two products. Purification by chromatography on silica gel (10:1 hexanes/ethyl acetate) afforded 5 g pure methyl 2-(4-iodophenyl)-2-methyl propanoate and 10 g mixture of the desired ester and 2-(4-iodophenyl)-2-methylpropanonitrile. ^1H NMR for methyl 2-(4-iodophenyl)-2-methyl propanoate (CDCl_3) δ 7.65 (d, $J = 8.5$, 2H), 7.09 (d, $J = 8.8$ Hz, 2H), 3.64 (s, 3H), 1.54 (s, 6H) ppm.

30

Part C. To 8 g of the crude mixture from Part B in THF (75 mL) and H_2O (25 mL) was added LiOH (3 g), and the reaction

was stirred overnight. Acid/base extraction afforded 3.6 g of 2-(4-iodophenyl)2-methylpropionic acid, Mass Spec (M+H)⁺ 290.8 and 5.3 g of 2-(4-iodophenyl)2-methylpropionitrile. IR(KBr) CN at 2236.66.

5

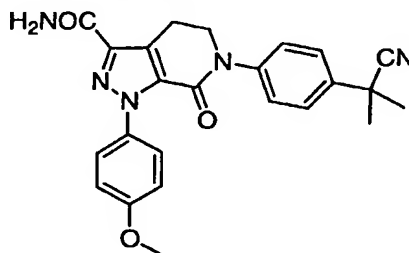
Part D. To a DMSO (4 mL, degassed) solution of ethyl 1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (0.6 g, 1.9 mmol), and 2-(4-iodophenyl) 2-methylpropionitrile (0.6 g, 2.2 mmol), and K₂CO₃ (0.66 g, 4.8 mmol) and was added CuI (73 mg, 0.3 mmol). The reaction was heated to 130°C for 18h. The reaction was cooled, extracted with EtOAc, washed with H₂O, and dried (MgSO₄). Purification by chromatography on silica gel (1:1 hexanes/ethyl acetate) afforded the title compound 0.4 g (45.9%) of a pale yellow solid; Mass Spec (M+H)⁺ 459.3.

15

Example 97

6-[4-(1-cyano-1-methylethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide

20



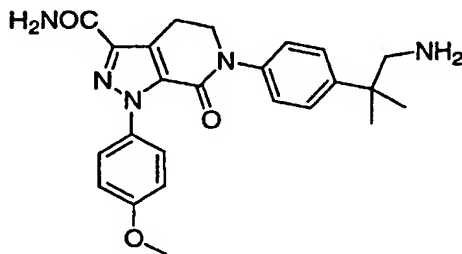
Ethyl 6-[4-(cyano-dimethyl-methyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (0.38 g, 0.83 mmol) obtained in Example 96 was placed in a sealed tube containing 10% ammonia in ethylene glycol (3 mL) and heated 80°C for 2h. The reaction was cooled, quenched with H₂O, extracted with EtOAc, and dried (MgSO₄). Recrystallization from

30

CH₂Cl₂/Hexanes afforded 0.31g (88%) of the title amide.
 High Resolution Mass Spec (M+H)⁺ for C₂₄H₂₄N₅O₃ 430.1898.

Example 98

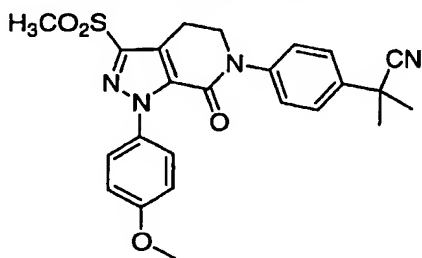
5 **6-[4-(2-Amino-1,1-dimethylethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide**



10 6-[4-(1-Cyano-1-methylethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide (0.1 g) was hydrogenated at 40psi in EtOH/HCl with 20 mg 10%Pd/C and purified by HPLC to afford 70 mg (56%) of title amine. High Resolution Mass Spec (M+H)⁺ for
 15 C₂₄H₂₈N₅O₃ 434.2176.

Example 99

1-(4-[(1-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridinyl-6-yl]phenyl)-2-methylpropanenitrile
 20

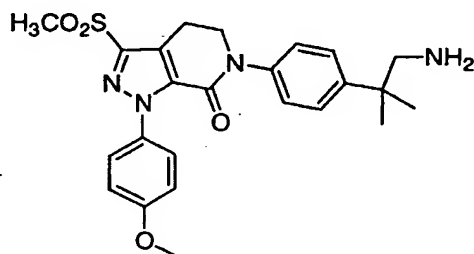


To a degassed DMSO (4 mL) solution containing 1-(4-methoxyphenyl)-3-methylsulfonyl-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.6 g, 1.8 mmol) and 2-(4-iodophenyl)-2-methylpropanenitrile (0.6 g, 2.2 mmol) was
 25

added K_2CO_3 (0.64 g, 4.6 mmol) and CuI (71 mg, 0.3 mmol). The reaction was heated to $130^\circ C$ for 18h. The reaction was cooled, extracted with EtOAc, washed with H_2O , and dried ($MgSO_4$). Purification by chromatography on silica gel (1:1
5 hexanes/ethyl acetate) afforded 0.52 g (61%) of a pale yellow foam; High Resolution Mass Spec $(M+H)^+$ for $C_{24}H_{25}N_4O_4S$ 456.1624.

Example 100

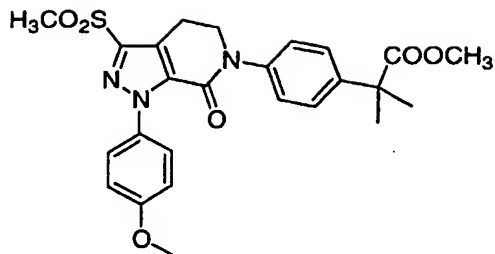
10 6-[4-(2-amino-1,1-dimethylphenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



15 1-{1-[(4-Methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridinyl-6-yl]phenyl}-2-methylpropanenitrile (0.1 g) was hydrogenated at 40psi in EtOH/HCl with 20 mg 10%Pd/C and purified by HPLC to afford 85 mg (68%) of the title amine. High Resolution Mass Spec
20 $(M+H)^+$ for $C_{24}H_{28}N_4O_4S$ 469.1907.

Example 101

Preparation of 2-{4-[3-methanesulfonyl-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydropyrazolo[3,4-c]pyridin-6-yl]phenyl}-2-methylproprionic acid methyl ester
25

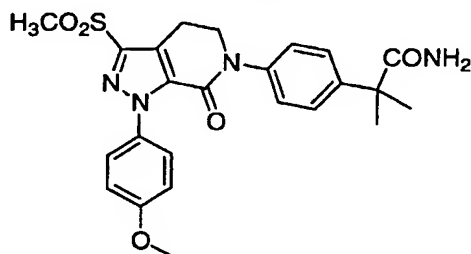


To a degassed DMSO (4 mL) solution was added ethyl 1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (0.4 g, 1.2 mmol) and methyl 2-(4-iodophenyl)-2-methyl propanoate (0.53 g, 1.7 mmol) was added K₂CO₃ (0.43 g, 3.1 mmol) and CuI (47 mg, 0.25 mmol). The reaction was heated to 130°C for 18h, cooled, extracted with EtOAc, washed with H₂O, and dried (MgSO₄). Purification by chromatography on silica gel (1:1 hexanes/ethyl acetate) afforded the titled compound 0.43 g (45.9%) of a pale yellow foam. High Resolution Mass Spec (M+H)⁺ for C₂₅H₂₈N₃O₆S 498.1691.

15

Example 102

2-(4-[1-(4-methoxyphenyl)-3-(methanesulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-2-methylpropanamide



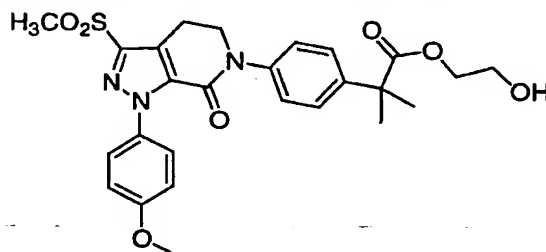
2-(4-[3-Methanesulfonyl-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydropyrazolo[3,4-c]pyridin-6-yl]phenyl)-2-methylproprionic acid methyl ester (0.095 g, 0.19 mmol) was placed in a sealed tube containing 10% ammonia in ethylene glycol (3 mL) and heated 80°C for 18h. The reaction was cooled, quenched with H₂O, extracted with EtOAc, and dried

(MgSO₄). Purification by HPLC afforded 35 mg (36%) title compound; High Resolution Mass Spec (M+H)⁺ for C₂₄H₂₇N₄O₅S 483.1725.

5

Example 103

2-Hydroxyethyl-2-(4-[1-(4-methoxyphenyl)-3-(methanesulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-2-methylpropanoate



10

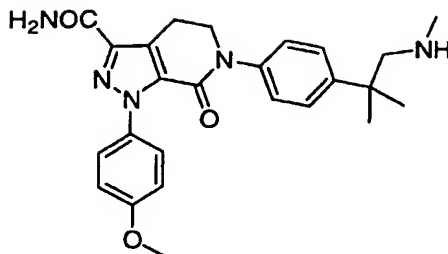
2-(4-[3-Methanesulfonyl-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydropyrazolo[3,4-c]pyridin-6-yl]phenyl)-2-methylproprionic acid methyl ester (0.077 g, 0.15 mmol) was placed in a sealed tube containing 10% ammonia in ethylene glycol (3 mL) and heated 80°C for 2h. The reaction was cooled, quenched with H₂O, extracted with EtOAc, and dried (MgSO₄). Purification by chromatography on silica (1:1 hexanes/ethyl acetate) and then HPLC purification afforded 27 mg (33%) title compound. High Resolution Mass Spec (M+H)⁺ for C₂₆H₃₀N₃O₇S 528.1776.

20

Example 104

6-(4-[1,1-dimethyl-2-(methylamino)ethyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide

25



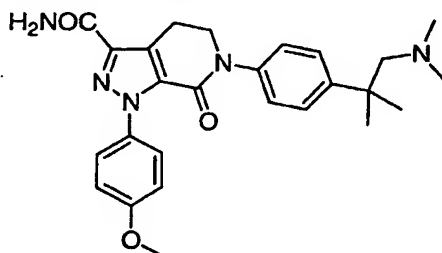
Part A. To ethyl 1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylate (2.35 g, 7.5 mmol) and 2-(4-iodophenyl)2-methylpropionic acid (2.6 g, 8.9 mmol) was added K_2CO_3 (3.1 g, 0.022 mol), DMSO (4 mL), and CuI (0.28 mg, 1.4 mmol). The reaction was heated to 130°C for 18h cooled, extracted with EtOAc, washed with H_2O , and dried ($MgSO_4$). Purification by chromatography on silica gel (5%MeOH/ CH_2Cl_2) afforded 1.1 g product.

Part B. To the acid from Part A (1 g, 2 mmol) in THF (30 mL) at 0°C was added 1M Borane in THF (2.5 mL, 2.5 mmol) and the reaction was allowed to stir 18h. The reaction was extracted with EtOAc, washed with brine, and dried (Na_2SO_4) to afford crude alcohol. To the alcohol was added CH_2Cl_2 (100 mL), molecular sieves, sodium acetate (0.17 g, 2 mmol), and pyridinium chlorochromate (0.72 g, 3.3 mmol) and the reaction was stirred 24h. After dilution with Et_2O , filtration through paper, and concentration, the crude residue was purified by chromatography on silica gel (2:1 hexanes/EtOAc) to afford 0.527 g of the desired aldehyde. 1H NMR ($CDCl_3$) δ 9.46 (s, 1H), 7.48 (d, J = 9.2 Hz, 2H), 7.35 (d, J = 8.7 Hz, 2H), 7.28 (d, J = 9.1 Hz, 2H), 6.92 (d, J = 9.2 Hz, 2H), 4.49 (q, J = 7 Hz, 2H), 4.13 (t, J = 6.6 Hz, 2H), 3.80 (s, 3H), 3.35 (t, J = 6.6 Hz, 2H), 1.45 (t, J = 7.3 Hz, 3H) ppm.

- Part C. To the aldehyde from Part B (95 mg, 0.2 mmol) in 1:1 THF/MeOH (5 mL) was added excess 33% methylamine in EtOH (0.1 mL). After 15 min 0.5M ZnCl₂ in THF (0.2 mL, 0.1 mmol) followed by sodium cyanoborohydride (13 mg, 0.2 mmol) were added. The reaction was stirred 24h. The solvents were removed and the residue was partitioned between EtOAc and H₂O. Extraction with EtOAc and drying (MgSO₄) afforded crude ester/amine.
- Part D. The ester/amine from Part C was heated in a sealed tube containing 2 mL of 10% NH₃/ethylene glycol at 80°C for 2h. After cooling the product was extracted by EtOAc, washed with water and dried (MgSO₄). Purification by HPLC and freeze-drying afforded the titled compound 78 mg (69%) as a white solid. High Resolution Mass Spec (M+H)⁺ for C₂₅H₃₀N₅O₃ 448.2337.

Example 105

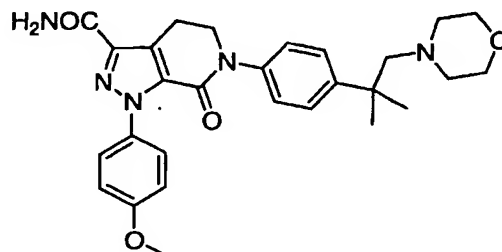
- 6-(4-[2-dimethylamino)-1,1-dimethylethyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



- Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₆H₃₂N₅O₃ 462.2529.

Example 106

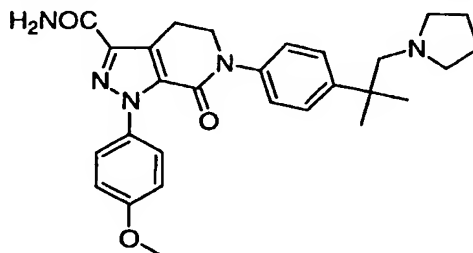
6-{4-[1,1-dimethyl-2-(1-morpholinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



5 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₄N₅O₄ 504.2637.

Example 107

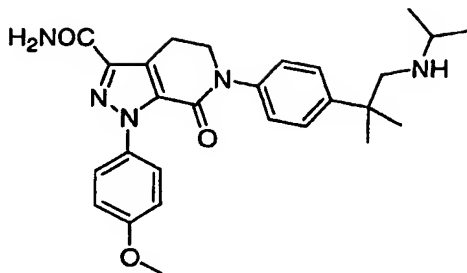
10 6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



15 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₄N₅O₃ 488.2667.

Example 108

20 6-{4-[2-(isopropylamino)1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide

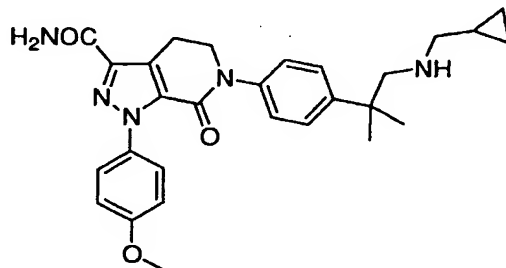


Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₇H₃₄N₅O₃ 476.2666.

5

Example 109

6-(4-{2-[(cyclopropylmethyl)amino]-1,1-dimethylethyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



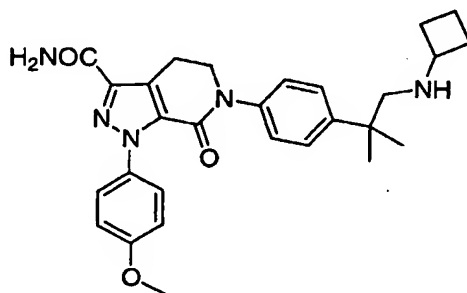
10

Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₃N₅O₃ 488.2670.

15

Example 110

6-{4-[2-(cyclobutylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide

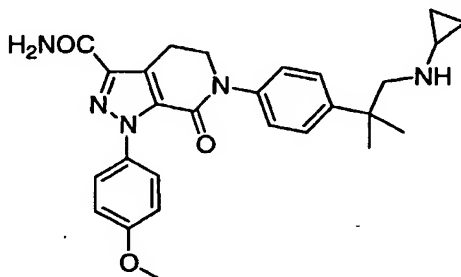


Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₄N₅O₃ 488.2668.

5

Example 111

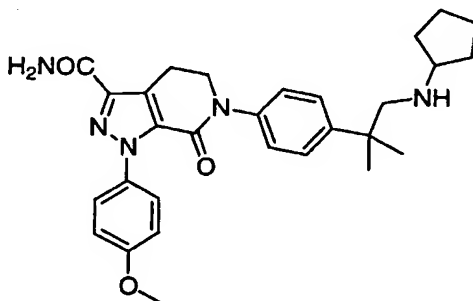
6-{4-[2-(cyclopropylamino)1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



10 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₇H₃₂N₅O₃ 474.2513

Example 112

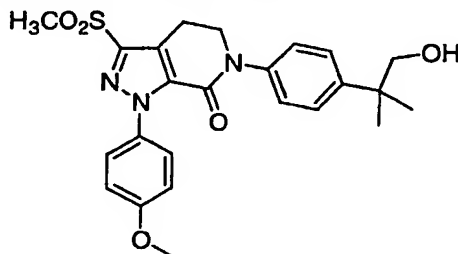
15 **6-{4-[2-(cyclopentylamino)1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide**



20 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₉H₃₆N₅O₃ 502.2814.

Example 113

6-[4-(2-hydroxy-1,1-dimethylethyl)phenyl]-1-(4-methoxyphenyl)-3-(methanesulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



5

Part A. To crude 2-{4-[3-methanesulfonyl-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydropyrazolo[3,4-c]pyridin-6-yl]phenyl}-2-methylproprionic acid methyl ester (2 g, 0.4 mmol) was added LiOH (0.5 g, 12 mmol) in THF/MeOH/H₂O for 24h. The reaction was acidified with 1N HCl and extracted with EtOAc and concentrated to afford crude acid as a semi solid mass.

Part B. The crude acid from Part A was then reduced with 1M borane in THF (7.3 mL, 7.3 mmol) in THF (25 mL) over 24h. The reaction was quenched with water and extracted with EtOAc and dried (MgSO₄) to afford the corresponding alcohol.

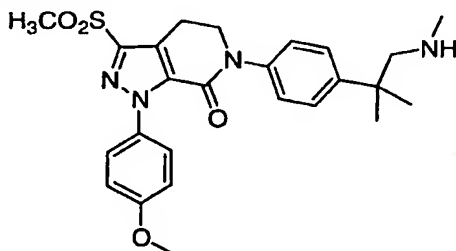
Part C. The crude alcohol from Part B (2.3 g, 4.9 mmol) was oxidized with pyridinium chlorochromate (1.7 g, 7.8 mmol), sodium acetate (0.4 g, 4.9 mmol), and molecular sieves in CH₂Cl₂ for 24h. Dilution with diethyl ether and filtration followed by chromatography on silica gel (1:1 hexanes/EtOAc) afforded 0.6 g (27%) of aldehyde; ¹H NMR CDCl₃ δ 9.46 (s, 1H), 7.48 (d, J = 8.8 Hz, 2H), 7.31 (m, 4H), 6.94 (d, J = 8.8 Hz, 2H), 4.15 (t, J = 6.6 Hz, 2H), 3.81 (s, 3H), 3.36 (t, J = 6.6 Hz, 2H), 3.31 (s, 3H), 1.44 (s, 6H) ppm.

30

Part D. To the aldehyde from Part C (34 mg, 0.072 mmol) was added 2-aminoimidazole sulfate (19 mg, 0.144 mmol) in 1:1 THF/MeOH (5 mL) followed by 0.5M ZnCl₂ (0.05 mL, 0.027 mmol) and 1M sodium cyanoborohydride in THF (0.07 mL, 0.07 mmol) and the reaction was stirred 24h. The reaction was quenched with water, extracted with EtOAc, and dried (MgSO₄). Purification by HPLC and freeze-drying afforded 12 mg (35%) of the desired alcohol: ¹H NMR (CDCl₃) δ 7.48 (d, J = 9.2 Hz, 2H), 7.41 (d, J = 8.4 Hz, 2H), 7.28 (d, J = 8.8 Hz, 2H), 6.94 (d, J = 9.1 Hz, 2H), 4.15 (t, J = 6.6 Hz, 2H), 3.81 (s, 3H), 3.61 (s, 2H), 3.34 (t, J = 6.6 Hz, 2H), 3.31 (s, 3H), 1.31 (s, 6H)ppm.

Example 114

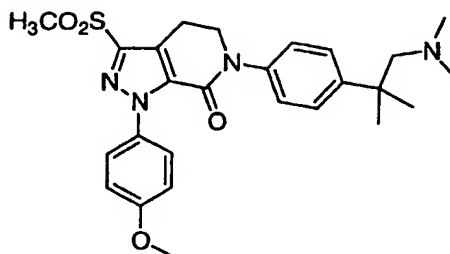
6-{4-[1,1-dimethyl-2-(methylamino)ethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₅H₃₁N₄O₄S 483.2049.

Example 115

6-{4-[2-(dimethylamino) 1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

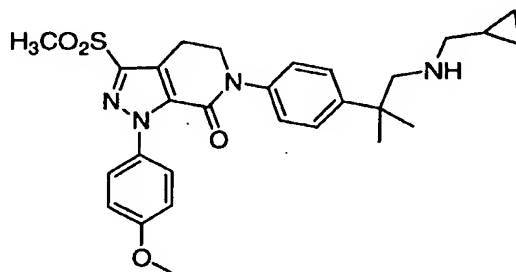


Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₆H₃₃N₄O₄S 497.2201.

5

Example 116

6-(4-{2-[(cyclopropylmethyl)amino]-1,1-dimethylethyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



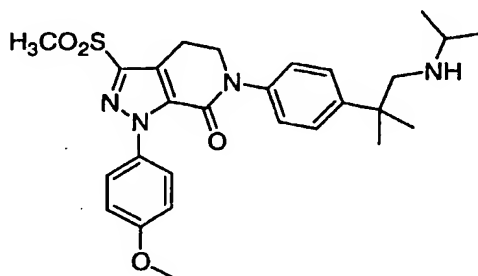
10

Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₅N₄O₄S 523.2362.

15

Example 117

6-{4-[1,1-dimethyl-2-(isopropylamino)ethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

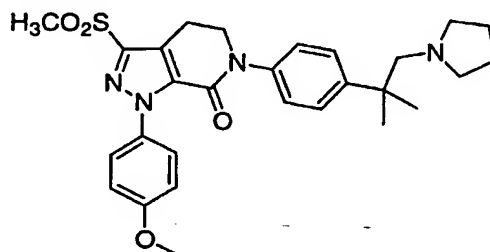


Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₇H₃₅N₄O₄S 511.2379.

5

Example 118

6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

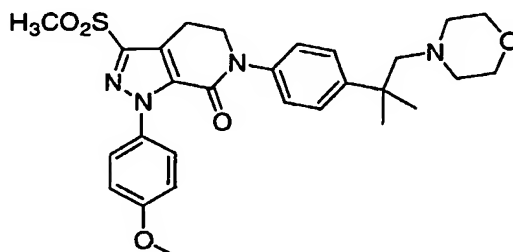


10 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₅N₄O₄S 523.2388.

15

Example 119

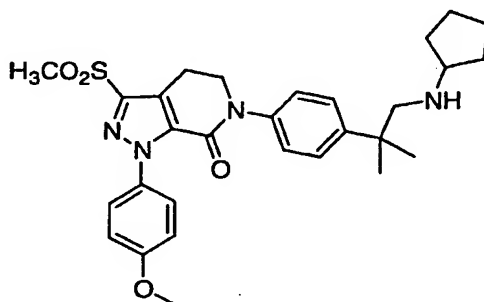
6-(4-[1,1-dimethyl-2-(1-morpholinyl)ethyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



20 Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₅N₄O₅S 539.2342.

Example 120

6-(4-[2-(cyclopentylamino) 1,1-dimethylethyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



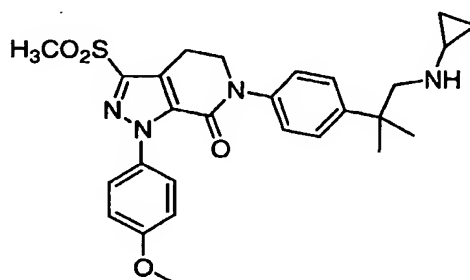
5

Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₉H₃₇N₄O₄S 537.2539.

10

Example 121

6-(4-{2-((cyclopropylamino) 1,1-dimethylethyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



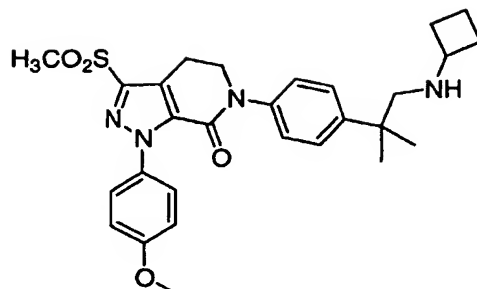
15

Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₇H₃₃N₄O₄S 509.2227.

20

Example 122

6-(4-[2-(cyclobutylamino) 1,1-dimethylethyl]phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

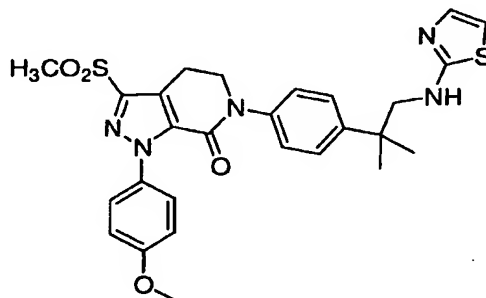


Following a procedure analogous to that used in Example 104, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₈H₃₅N₄O₄S 523.238.

5

Example 123

6-{4-[1,1-dimethyl-2-(1,3-thiazol-2-yl amino)ethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,7-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one

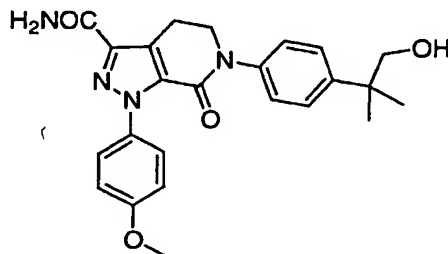


10

Prepared as previously described above. High Resolution Mass Spec (M+H)⁺ for C₂₇H₃₀N₅O₄S₂ 552.1727.

Example 124

6-[4-(2-hydroxy-1,1-dimethylethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide

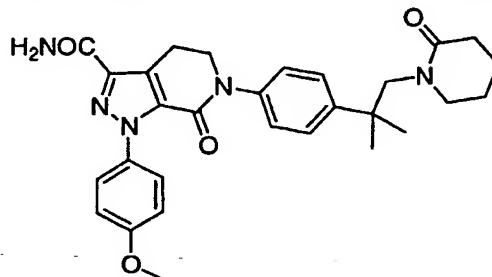


Following a procedure analogous to that used in Example 113, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₄H₂₇N₄O₄ 435.2016.

5

Example 125

6-{4-[1,1-dimethyl-2-(2-oxo-1-piperidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide



10

To 6-[4-(1-cyano-1-methylethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide (0.17 g, 0.39 mmol) was hydrogenated at 40psi in ethanol with 0.5 mL conc. HCl and 10% palladium on carbon (25 mg) for 72h. The reaction was filtered and concentrated. To the amine in THF (5 mL) at 0°C was added 5-bromovaleryl chloride (99 mg, 0.5 mmol) and TEA (1 mL) and the reaction was stirred 1h. To the reaction was added potassium *t*-butoxide (0.24 g, 1.9 mmol) and the reaction was stirred 24h. The reaction was quenched with water and extracted with ethyl acetate and dried (MgSO₄). Purification by HPLC and freeze-drying afforded 15 mg (7.5%). Mass Spec (M+H)⁺ 516.3.

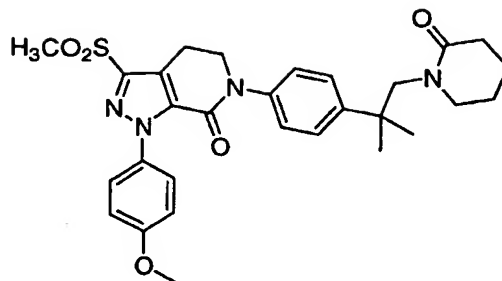
15

20

25

Example 126

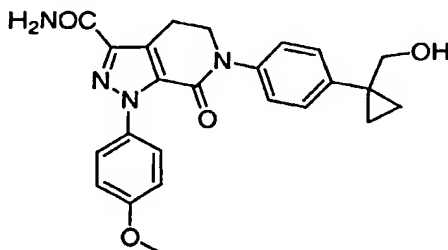
6-{4-[1,1-dimethyl-2-(2-oxo-1-piperidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one



1- $\{4-[(1\text{-Methoxyphenyl})-3\text{-(methoxysulfonyl)}-7\text{-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridinyl-6-yl}]$
 phenyl}-2-methylpropanenitrile was converted into the
 5 target compound by the same procedure as that of Example
 125. Mass Spec (M+H)⁺ 551.3.

Example 127

10 6-[4-(1-Hydroxymethylcyclopropyl)phenyl]-1-(4-methoxy-
 phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
 c]pyridine-3-carboxylic acid amide

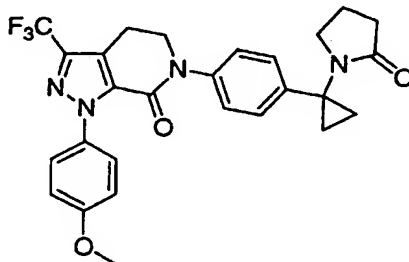


Following a procedure analogous to that used in Example 95,
 the title compound was prepared. LC/MS (ESI⁺) 433.4 (M+H)⁺.

15

Example 128

1-(4-Methoxyphenyl)-6-{4-[1-(2-oxo-pyrrolidin-1-yl)-
 cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-
 pyrazolo[3,4-c]pyridin-7-one



20

Part A. 4-Iodophenylcyclopropyl carboxylic acid (7.42 g, 25.76 mmol) was stirred in CH₂Cl₂ (60 mL) at rt under N₂. Et₃N (5.4 mL, 38.64 mmol, 1.5 eq) was added followed by the addition of DPPA (8.27 mL, 38.64 mmol, 1.5 eq). The resulting mixture was stirred at rt overnight. It was poured into ice H₂O (100 mL), acidified with 6N HCl, and then extracted with CH₂Cl₂. The organic layer was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated under vacuum. The oil residue obtained was dissolved in t-BuOH (40 mL) and refluxed for 2-3 h. After cooling, the solvent was evaporated. The residue was purified by FCC (silica gel, hexane: CH₂Cl₂=1:1, then CH₂Cl₂, then CH₂Cl₂:MeOH=100:1 to 25:1) to give pure [1-(4-iodophenyl)-cyclopropyl]-carbamic acid tert-butyl ester (6.01 g, yield: 65%). This compound (2.12 g, 5.89 mmol) was stirred in CH₂Cl₂ (10 mL) and TFA (10 mL) at rt for 2h. After evaluation, the residue was taken up in CHCl₃ (100 mL) and H₂O (100 mL). The aqueous layer was basified with K₂CO₃, extracted with CHCl₃ (2 x), dried over MgSO₄, filtered, and concentrated to dryness to give pure 1-(4-iodophenyl)cyclopropylamine (1.50 g, yield: 98%). ¹H NMR (CDCl₃) δ 7.62 (m, 2H), 7.06 (m, 2H), 1.88 (m, 2H), 1.07 (m, 2H), 0.95 (m, 2H) ppm. HRMS C₉H₁₁IN (M+H)⁺ 259.9930 calcd for 259.9936.

25

Part B. The mixture of the product from Part A (0.32 g, 1.24 mmol), NaOH (0.15 g, 3.72 mmol, 3.0 eq), and 4-chlorobutyryl chloride (0.18 mL, 1.61 mmol, 1.3 eq) was stirred in CH₂Cl₂ (7 mL) at rt for 1 h under N₂. H₂O was added. It was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, and concentrated to dryness. The residue was dissolved in THF (10 mL). KOtBu (0.40 g, 4.16 mmol) was added as one portion. The mixture was

30

stirred at 0°C under N₂ for 30 min. EtOAc was added. It was washed with H₂O and brine, dried over MgSO₄, and concentrated to produce 1-[1-(4-iodophenyl)cyclopropyl]-pyrrolidin-2-one (0.27 g, yield: 100%). ¹H NMR (CDCl₃) δ

5 7.68 (d, J=8.5 Hz, 2H), 6.87 (d, J=8.5 Hz, 2H), 3.26 (t, J=7.3 Hz, 2H), 2.26 (t, J=7.6 Hz, 2H) 1.88 (t, J=7.0 Hz, 2H), 1.22 (m, 2H), 1.10 (m, 2H) ppm.

Part C. The product from Part B (64 mg, 0.196 mmol) and 1-

10 (4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.061 g, 0.196 mmol) were stirred in DMSO (0.3 mL) under N₂. K₂CO₃ (0.067 g, 0.49 mmol, 2.5 eq) was added, followed by the addition of CuI (0.037 g, 0.194 mmol) and 1,10-phenanthroline (0.020 g,

15 0.108 mmol). The resulting mixture was heated at 120°C for 3h. After cooling, it was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, CH₂Cl₂:EtOAc=1:1, then EtOAc) to give 1-(4-

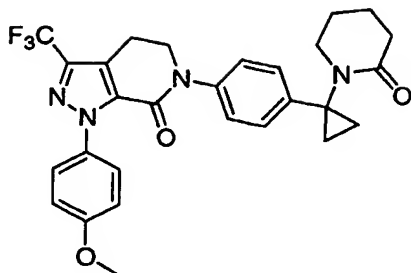
20 methoxyphenyl)-6-{4-[1-(2-oxo-pyrrolidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one (45 mg, yield: 45%). ¹H NMR (CDCl₃) δ 7.46 (d, J=8.8 Hz, 2H), 7.25 (AA'BB', J=8.6 Hz, 4H), 6.91 (d, J=9.2 Hz, 2H), 4.10 (t, J=6.6 Hz, 2H), 3.81

25 (s, 3H), 3.38 (t, J=7.2 Hz, 2H), 3.14 (t, J=6.6 Hz, 2H), 2.37 (t, J=7.7 Hz, 2H), 1.98 (q, J=7.7 Hz, 2H), 1.32 (m, 2H), 1.21 (m, 2H) ppm. HRMS C₂₇H₂₆F₃N₃O₄ (M+H)⁺ 511.1931 calcd for 511.1958.

30

Example 129

1-(4-Methoxyphenyl)-6-{4-[1-(2-oxo-piperidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Following the procedures analogous to those used in Example 128, the title compound was prepared. The product was purified by RP-prep LC-MS (35-98% CH₃CN/H₂O in a 10-min

5 run). HRMS C₂₉H₃₂O₂F₃N₄ (M+H)⁺ 525.2486 calcd for 525.2477.

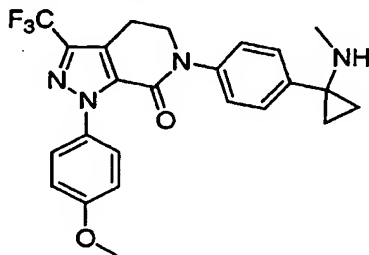
¹H NMR (CDCl₃) δ 7.46 (d, J=8.8 Hz, 2H), 7.25 (d, J=8.4 Hz, 2H), 6.91 (d, J=9.2 Hz, 2H), 4.10 (t, J=6.6 Hz, 2H), 3.82 (s, 3H), 3.37 (t, J=6 Hz, 2H), 3.15 (t, J=6.6 Hz, 2H), 2.51 (t, J=6 Hz, 2H), 1.79 (m, 4H), 1.33 (m, 2H), 1.29 (m, 2H)

10 ppm.

Example 130

1-(4-Methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)-phenyl]-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one

15



Part A. 1-(4-Iodophenyl)cyclopropyl-carbamic acid tert-butyl ester (2.14 g, 5.85 mmol) was stirred in THF (20 mL) at 0°C under N₂. MeI (3 mL) was added followed by

20 portionwise addition of NaH (2.34 g, 5 eq). The reaction was stirred at rt overnight. Several drops of H₂O and EtOAc (20 mL) were added to quench the reaction. The organic solvent was evaporated, and H₂O was added. It was extracted with Et₂O (2x), washed with brine, dried over

MgSO₄, and concentrated to dryness. The residue was purified by FCC (silica gel, CH₂Cl₂:hexanes=0:1 to 1:1 to 1:0) to give pure [1-(4-iodophenyl)cyclopropyl]-methyl-carbamic acid tert-butyl ester as a white solid (2.07 g, 5 yield 95%). LC/MS (ESI⁺) 373.8 (M+H), t_R=2.95 min (10-90% CH₃CN in H₂O in a 4-min run).

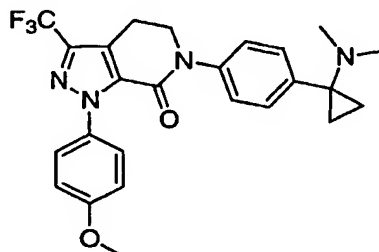
Part B. The product from Part A (205 mg, 0.55 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-10 7H-pyrazolo[3,4-c]pyridin-7-one (170 mg, 0.55 mmol) were stirred in DMSO (0.4 mL) under N₂. K₂CO₃ (250 mg, 1.81 mmol, 3.3 eq) was added, followed by the addition of CuI (52 mg, 0.27 mmol, 0.5 eq) and 1,10-phenanthroline (50 mg, 0.27 mmol, 0.5 eq). The resulting mixture was heated at 15 120°C for 2h. After cooling, it was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by flash column chromatography (silica gel, CH₂Cl₂:EtOAc=1:1, then EtOAc) to give (1-{4-[1-(4-methoxyphenyl)-7-oxo-3-20 trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-methyl-carbamic acid tert-butyl ester (250 mg, yield: 82%). ¹H NMR (CDCl₃) δ 7.46 (d, J=9.2 Hz, 2H), 7.23 (d, J=8.4 Hz, 2H), 7.09 (m, 2H), 6.91 (d, J=9.1 Hz, 2H), 4.12 (t, J=6.6 Hz, 2H), 3.81 (s, 3H), 3.15 25 (t, J=6.6 Hz, 2H), 2.90 (s, br, 3H), 1.42 (s, br, 9H), 1.33 (m, 2H), 1.20 (m, 2H) ppm. LC/MS (ESI⁺) 557.4.

Part C. The product from Part B (250 mg, 0.45 mmol) was stirred in CH₂Cl₂ (2 mL) and TFA (2 mL) at rt for 20 min. 30 The solvents were evaporated. The residue was purified by FCC (silica gel, EtOAc, then EtOAc: MeOH=10:1) to yield the title compound (188 mg, 92%). ¹H NMR (CDCl₃) δ 7.55 (d, J=8.4 Hz, 2H), 7.45 (d, J=9.2 Hz, 4H), 7.36 (d, J=8.4 Hz, 2H), 6.91 (d, J=8.8 Hz, 2H), 4.15 (t, J=6.6 Hz, 2H), 3.80

(s, 3H), 3.17 (t, $J=6.6$ Hz, 2H), 2.50 (s, 3H), 1.56 (m, 2H), 1.12 (m, 2H) ppm. LC/MS (ESI⁺) 457.4.

Example 131

5 **6-[4-(1-Dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one**

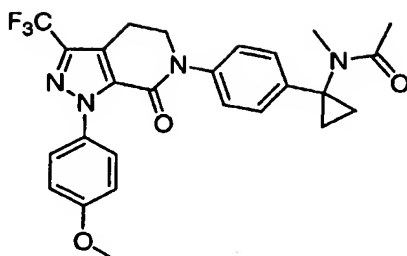


The product from Example 130 (30 mg, 0.066 mmol) was stirred in CH₃CN (0.2 mL) at rt under N₂. Aqueous formaldehyde (0.07 mL, 7 mmol, 10 eq) was added followed by the addition of HOAc (0.012 mL, 0.21 mmol, 3.2 eq). The mixture was stirred for 15 min, and then NaBH₃CN (12 mg, 0.198 mmol) was added. The mixture was stirred at rt for 2h. Several drops of acetone were added followed by 1N NaOH. The mixture was extracted with CH₂Cl₂, washed with H₂O and brine, dried over MgSO₄, and concentrated to dryness. The residue was purified by FCC (silica gel, EtOAc, than EtOAc: MeOH=10:1) to yield the title compound (15.7 mg, 51%). ¹H NMR (CDCl₃) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.29 (m, 4H), 6.92 (d, $J=9.2$ Hz, 2H), 4.15 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.16 (t, $J=6.6$ Hz, 2H), 2.28 (s, 6H), 1.02 (m, 2H), 0.81 (m, 2H) ppm. LC/MS (ESI⁺) 471.4.

25

Example 132

N-(1-(4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-cyclopropyl)-N-methyl-acetamide



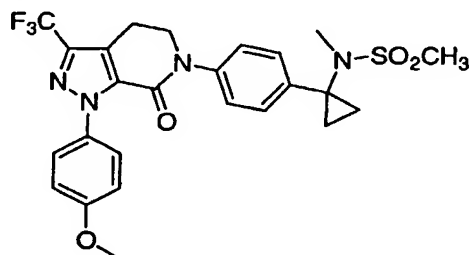
Following a procedure analogous to that used for the preparation of Example 93, the title compound was prepared. Silica gel purification yielded the title compound. LC/MS

5 (ESI⁺) 499.4 (M+H). ¹H NMR (CDCl₃) δ 7.46 (d, J=9.2 Hz, 2H), 7.26 (m, 2H), 6.93 (AA'BB', J=8.8, 7.0 Hz, 4H), 4.12 (t, J=6.6 Hz, 2H), 3.81 (s, 3H), 3.16 (t, J=6.6 Hz, 2H), 3.01 (s, 3H), 2.05 (s, 3H), 1.50 (m, 4H) ppm.

10

Example 133

***N*-(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-methanesulfonamide**

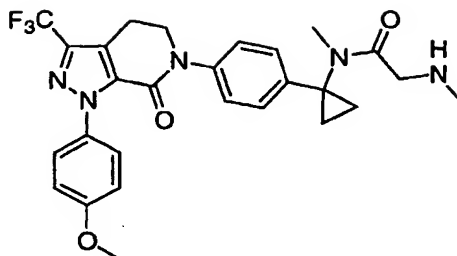


15 Following a procedure analogous to that used for the preparation of Example 94, the title compound was prepared. Silica gel purification yielded the pure desired product. LC/MS (ESI⁺) 535.6 (M+H)⁺.

20

Example 134

***N*-(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-2-methylaminoacetamide**

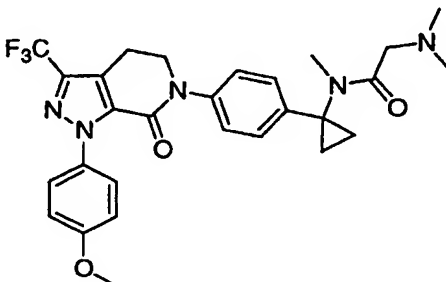


Part A. The product of Example 130 (45 mg, 0.1 mmol) was stirred in CH_2Cl_2 (1 mL) at rt. NaOH (12 mg, 3.0 eq) was added followed by the addition of chloroacetyl chloride (0.015 mL, 2.0 eq). The mixture was stirred at rt for 3h. Additional NaOH (20 mg) and chloroacetyl chloride (0.020 mL) were added. The mixture was stirred at rt overnight. The mixture was extracted with CH_2Cl_2 (2x), washed with H_2O and brine, dried over MgSO_4 , filtered, and concentrated to dryness. The residue was used directly in the next step without further purification. LC/MS (ESI⁺) 533.6 (M+H), t_R =2.63 min (10-90% CH_3CN in H_2O in a 4-min run).

Part B. The product from part A (15 mg, 0.028 mmol) was stirred in DMF (0.1 mL) in a Pyrex tube under N_2 . K_2CO_3 (20 mg) was added, followed by the addition of a solution of NHMe_2 in THF (2M, 0.1 mL). The reaction mixture was stirred at 80°C overnight. H_2O was added, and the mixture was extracted with EtOAc, washed with brine, dried over MgSO_4 , filtered, and concentrated. The residue was purified by FCC (silica gel, CH_2Cl_2 , then CH_2Cl_2 :EtOAc, then EtOAc:MeOH=10:1) to give the title compound (5.0 mg, yield: 33%). ^1H NMR (CDCl_3) δ 7.46 (d, J =8.8 Hz, 2H), 7.29 (m, 4H), 6.92 (d, J =9.2 Hz, 2H), 4.15 (t, J =6.6 Hz, 2H), 3.81 (s, 3H), 3.16 (t, J =6.6 Hz, 2H), 2.28 (s, 6H), 1.02 (m, 2H), 0.81 (m, 2H) ppm. LC/MS (ESI⁺) 528.6 (M+H), t_R =2.07 min (10-90% CH_3CN in H_2O in a 4-min run).

Example 135

2-Dimethylamino-N-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N-methylacetamide



5

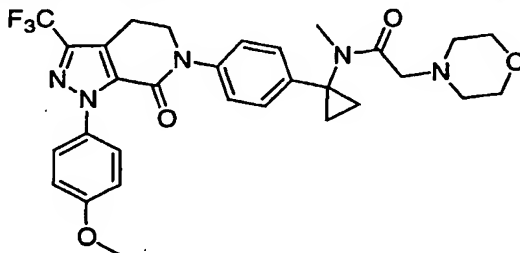
Following a procedure analogous to that used for the preparation of Example 134, the title compound was prepared. Silica gel purification yielded the pure desired product. LC/MS (ESI⁺) 542.6 (M+H), t_R=2.10 min (10-90%

10 CH₃CN in H₂O in a 4-min run).

Example 136

N-(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-cyclopropyl)-N-methyl-2-morpholin-4-yl-acetamide

15

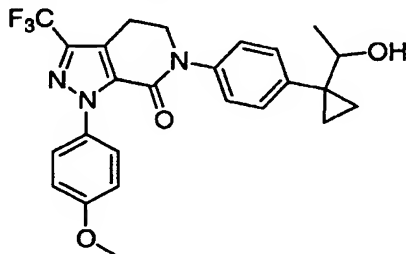


Following a procedure analogous to that used for the preparation of Example 134, the title compound was prepared. Silica gel purification yielded the pure desired

20 product. LC/MS (ESI⁺) 584.2 (M+H)⁺, t_R=2.05 min (10-90% CH₃CN/H₂O in a 4-min run).

Example 137

6-{4-[1-(1-Hydroxyethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



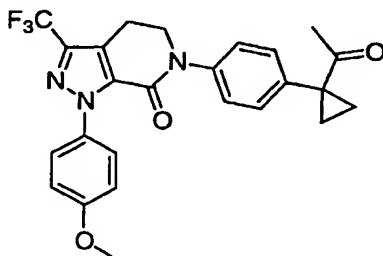
5

1-(4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropanecarbaldehyde (93 mg, 0.21 mmol) was stirred in Et₂O (2 mL) at -78°C. ZnMe₂ (2M in toluene, 0.16 mL, 1.5 eq) was added followed by the addition of TiCl₄ (1 M in CH₂Cl₂, 0.3 mL). The resulting mixture was stirred for 1h. The reaction was quenched by addition of NH₄Cl, extracted with EtOAc, washed with H₂O and brine, dried over MgSO₄, filter, and concentrated. The residue was purified by silica gel column to yield the pure desired product (62 mg, yield: 64.5%). LC/MS (ESI⁺) 472.6 (M+H)⁺, t_R=2.44 min (35-95% CH₃CN/H₂O in a 6-min run).

15

Example 138

20 6-[4-(1-Acetylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



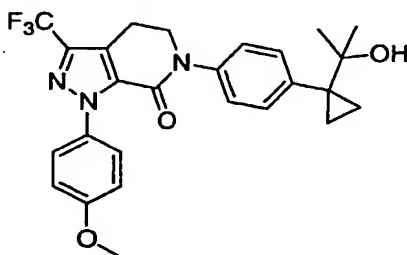
The product from Example 137 (30 mg, 0.063 mmol) was stirred in CH₂Cl₂ (1 mL) at rt under N₂. 4Å molecular sieves

25

(30 mg) and NaOAc (15.4 mg, 0.187 mmol) were added followed by the addition of PCC (27.5 mg, 0.126 mmol). The reaction mixture was stirred at rt for 1.5 h. The mixture was filtered through Celite®, rinsed with CH₂Cl₂, washed with H₂O, brine, concentrated to dryness. Silica gel purification afforded the title compound. LC/MS(ESI⁺) 470.6 (M+H)⁺, t_R=2.77 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (CDCl₃) δ 7.47 (d, J=8.4 Hz, 2H), 7.33 (AA'BB', J=8.8 Hz, 4H), 6.93 (dd, J=8.8, 2.3 Hz, 2H), 4.16 (t, J=6.6 Hz, 2H), 3.83 (s, 3H), 3.29 (t, J=6.6 Hz, 2H), 2.77 (s, 6H), 1.67 (m, 2H), 1.15 (m, 2H) ppm.

Example 139

6-{4-[1-(1-Hydroxy-1-methyl-ethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydropyrazolo[3,4-c]pyridin-7-one



Part A. 1-(4-Iodo-phenyl)-cyclopropanecarboxylic acid methyl ester (0.96 g, 3.17 mmol) was stirred in THF (15 mL) at -78°C under N₂. MeMgCl (3.0 M in THF, 4.2 mL, 4.0 eq) was added dropwise, and the reaction was stirred for 1 h during which period the temperature was raised from -78°C to 0°C. It was quenched by the addition of sat'd NH₄Cl, and extracted with EtOAc (2 x). The organics were washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, hexanes, then hexanes:CH₂Cl₂=1:1 to 0:1) to give 2-[1-(4-iodo-phenyl)-cyclopropyl]-propan-2-ol (0.71 g,

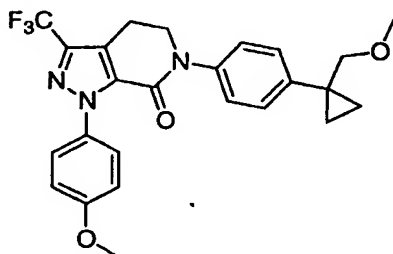
yield: 73.9%). LC/MS(ESI⁺) 303.4 (M+H)⁺, t_R=2.57 min (10-90% CH₃CN/H₂O in a 4-min run).

Part B. The product from Part A (102 mg, 0.33 mmol) and
5 (105 mg, 0.34 mmol) were stirred in dry DMSO (0.5 mL).
K₂CO₃ (90.5 mg, 2.0 eq) was added followed by the addition
of CuI (32 mg, 0.17 mmol) and 1,10-phenanthroline (31 mg,
0.17 mmol). The resulting mixture was heated at 120°C for
3h. After cooling, it was extracted with EtOAc (2x),
10 washed with H₂O and brine, dried over MgSO₄, filtered, and
concentrated to dryness. The residue was purified by FCC
(silica gel, CH₂Cl₂:EtOAc=1:1, then EtOAc) to give the title
compound (95 mg, yield: 59.4%). LC/MS(ESI⁺) 486.8 (M+H)⁺,
t_R=3.03 min (10-90% CH₃CN/H₂O in a 4-min run).

15

Example 140

6-[4-(1-Methoxymethylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydropyrazolo[3,4-c]pyridin-7-one



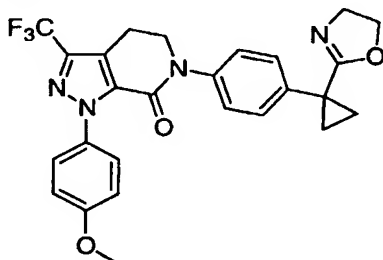
20

Part A. [1-(4-Iodo-phenyl)-cyclopropyl]-methanol (0.25 g,
0.94 mmol) was dissolved in CH₂Cl₂ (1.5 mL). Proton sponge
(0.21 g, 0.97 mmol) was added followed by
trimethoxyloxonium tetrafluoroborate (0.15 g, 1.0 mmol).
25 The reaction was allowed to stir for 3 h and was then
quenched with H₂O, concentrated, and purified via flash
chromatography (silica, 100% EtOAc) to afford the title
compound (0.13 g, yield: 47%). ¹H NMR (CDCl₃) δ 7.64 (d,
J=8.4 Hz, 2H), 7.11 (d, J=8.2 Hz, 2H), 3.65 (d, J=6.2 Hz,
30 2H), 1.57 (s, 3H), 0.86 (s, 4H) ppm.

Part B. The product from Part A and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one were coupled using the usual Buchwald
 5 Ullman procedure. LC/MS (ESI⁺) 472.6 (M+H)⁺, t_R=2.98 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR ((CD₃)₂CO, 300 MHz) δ 7.49 (d, J=9.1 Hz, 2H), 7.29 (AA'BB', J=8.8 Hz, 4H), 6.96 (dd, J=9.2 Hz, 2H), 4.15 (t, J=6.6 Hz, 2H), 3.82 (s, 3H), 3.44 (s, 2H), 3.23 (s, 3H), 3.16 (t, J=6.3 Hz, 2H),
 10 0.84 (d, J=2.2 Hz, 2H), 0.81 (d, J=2.5 Hz, 2H) ppm.

Example 141

6-{4-[1-(4,5-Dihydro-oxazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-
 15 pyrazolo[3,4-c]pyridin-7-one



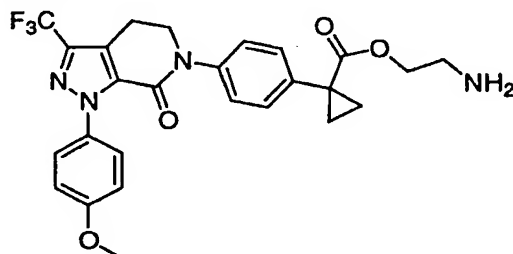
Part A. To a slurry of 1-(4-iodo-phenyl)-cyclopropane-carboxylic acid (0.693 g, 2.41 mmol) in CH₂Cl₂ (3.0 mL) at 0°C was added (COCl)₂ (0.40 mL, 4.6 mmol) dropwise. The
 20 reaction was warmed to rt and stirred under N₂ for 1 h. The reaction was monitored by LC/MS. Upon completion the reaction was concentrated on the rotary evaporator and diluted with CH₂Cl₂ (3 mL). Ethanolamine (0.30 mL, 4.54 mmol) was added drop-wise and the reaction stirred for 1.5
 25 h. The reaction was then quenched with H₂O and extracted with EtOAc (2x). The organic layers were washed with brine, dried over Na₂SO₄, filtered, and concentrated to dryness. The crude 1-(4-iodo-phenyl)-cyclopropanecarboxylic acid (2-hydroxy-ethyl)-amide was

taken directly to the next reaction without further purification. LC/MS (ESI⁺) 332.2 (M+H)⁺, t_R=2.16 min (10-90% CH₃CN/H₂O in a 4-min run). It was dissolved in THF (10.0 mL) and methoxycarbonylsulfamoyl triethylammonium hydroxide inner salt (0.61 g, 2.56 mmol) was added. The reaction was heated to 70°C for 2h and then cooled. The reaction mixture was diluted with EtOAc and washed with H₂O (2x), brine, dried over Na₂SO₄, filtered, and concentrated to dryness. The residue was purified by flash chromatography (silica, EtOAc: Hexanes 3:1) to 2-[1-(4-iodo-phenyl)-cyclopropyl]-4,5-dihydro-oxazole (0.41 g, yield: 55%). LC/MS (ESI⁺) 314.0 (M+H), t_R=1.62 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (CDCl₃) δ 7.65 (d, J=8 Hz, 2H), 7.13 (d, J=8.4 Hz, 2H), 4.32 (t, J=9.5 Hz, 2H), 3.87 (t, J=9.1, 9.6, 2H), 1.67 (m, 2H), 1.23 (m, 2H) ppm.

Part B. The product from Part A (75.2 mg, 0.240 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (76.8 mg, 0.247 mmol) were dissolved in DMSO (0.5 mL). Potassium carbonate (0.109 g, 0.788 mmol), copper iodide (spatula tip), and 1,10-phenanthroline (spatula tip) were added and the reaction was heated to 120°C for 12h under an environment of N₂. The reaction was cooled, diluted with EtOAc, washed with H₂O (2x), brine, dried over Na₂SO₄, filtered, and concentrated in vacuo. Flash chromatography (silica, 100% EtOAc) afforded the title compound. LC/MS (ESI⁺) 497.6 (M+H), t_R=2.44 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR ((CD₃)₂CO, 300 MHz) δ 7.51 (d, J=9.0 Hz, 2H), 7.32 (AA'BB', J=8.4 Hz, 4H), 6.97 (d, J=9.0 Hz, 2H), 4.16 (m, 4H), 3.82 (s, 3H), 3.67 (t, J=9.2 Hz, 2H), 3.17 (t, J=6.6 Hz, 2H), 2.02 (m, 2H), 1.42 (m, 2H), 1.17 (m, 2H) ppm.

Example 142

1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropanecarboxylic acid 2-amino-ethyl ester



5

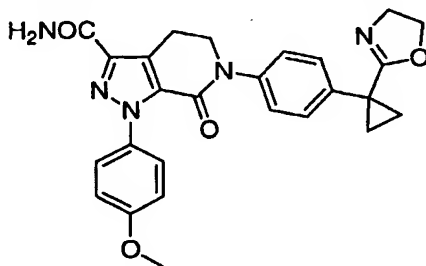
A side product resulting from a minor impurity in the starting material of Part C of Example 141 was isolated and characterized to be the title compound. LC/MS (ESI⁺) 515.6 (M+H)⁺, t_R=2.22 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H

10 NMR (CD₃)₂CO, δ 7.50 (d, J=8.8 Hz, 2H), 7.38 (AA'BB', J=8.6 Hz, 4H), 7.00 (d, J=8.8 Hz, 2H), 4.20 (t, 2H), 3.83 (s, 3H), 3.45 (t, 2H), 3.20 (m, 4H), 1.40 (m, 2H), 0.95 (m, 2H) ppm.

15

Example 143

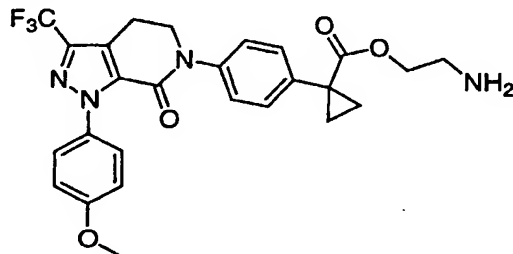
6-{4-[1-(4,5-Dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



20 Part A. The product of Part A from Example 141 (0.10 g, 0.32 mmol) and 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.10 g, 0.32 mmol) were dissolved in DMSO (1.5 mL). Potassium carbonate (1.3 g, 0.94 mmol), copper iodide
25 (0.02 g, 0.10 mmol), and 1,10-phenanthroline (0.02 g, 0.11

Example 142

1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropanecarboxylic acid 2-amino-ethyl ester



5

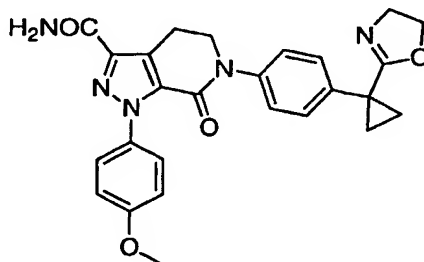
A side product resulting from a minor impurity in the starting material of Part C of Example 141 was isolated and characterized to be the title compound. LC/MS (ESI⁺) 515.6 (M+H)⁺, t_R=2.22 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H

10 NMR (CD₃)₂CO, δ 7.50 (d, J=8.8 Hz, 2H), 7.38 (AA'BB', J=8.6 Hz, 4H), 7.00 (d, J=8.8 Hz, 2H), 4.20, (t, 2H), 3.83 (s, 3H), 3.45 (t, 2H), 3.20 (m, 4H), 1.40 (m, 2H), 0.95 (m, 2H) ppm.

15

Example 143

6-{4-[1-(4,5-Dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



20 Part A. The product of Part A from Example 141 (0.10 g, 0.32 mmol) and 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.10 g, 0.32 mmol) were dissolved in DMSO (1.5 mL). Potassium carbonate (1.3 g, 0.94 mmol), copper iodide
25 (0.02 g, 0.10 mmol), and 1,10-phenanthroline (0.02 g, 0.11

mmol) were then added and the reaction was heated to 120°C for 12 h under an environment of N₂. The reaction was cooled, diluted with EtOAc, washed with H₂O (2x), brine, dried over Na₂SO₄, filtered, and concentrated to dryness to give 6-{4-[1-(4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester. LC/MS (ESI⁺) 501.8 (M+H)⁺, t_R=2.64 min (10-90% CH₃CN/H₂O in a 4-min run).

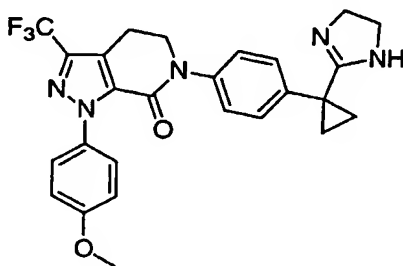
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Part B. The product from Part A (0.10 g, 0.20 mmol) was dissolved in ethylene glycol saturated with ammonia (2.0 mL) and heated to 85°C for 4 h. Reaction was cooled, diluted with H₂O, and washed with EtOAc (3x). Organic portions were combined and washed with brine (2x), dried over Na₂SO₄, filtered, and concentrated to dryness. The title compound was recrystallized from EtOAc/Hexanes (0.03 g, yield: 32%). LC/MS (ESI⁺) 472.6 (M+H)⁺, t_R=1.49 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (CD₃OD) δ 7.47 (d, J=9.2 Hz, 2H), 7.38 (AA'BB', J=8.5 Hz, 4H), 4.86 (t, 2H), 4.25 (t, 2H), 3.81 (s, 3H), 3.72 (t, 2H), 1.49 (m, 2H), 1.16 (m, 2H) ppm.

20

Example 144

25 6-{4-[1-(4,5-Dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Part A. 1-(4-Iodo-phenyl)-cyclopropanecarboxylic acid (0.99 g, 3.4 mmol), DCC (0.71 g, 3.4 mmol), and pentafluorophenol

30

(0.91 g, 4.9 mmol) were added into CH₂Cl₂ (6 mL) and allowed to stir for 2 h. Piperidine (0.7 mL, 7.1 mmol) was then added dropwise to the slurry. Reaction was allowed to stir for an additional 12 h. The reaction was then diluted with
5 EtOAc; filtered; washed with 1N HCl, 1N NaOH (2x), and brine; dried over MgSO₄; filtered; and concentrated to dryness. The crude mixture was purified by flash chromatography (silica, EtOAc:hexanes (3:1) to give [1-(4-iodo-phenyl)-cyclopropyl]-piperidin-1-yl-methanone (1.09,
10 yield: 88%). ¹H NMR (CDCl₃, 300 MHz) δ 7.60 (d, J=8.8 Hz, 2H), 6.92 (d, J=8.4 Hz, 2H), 3.55 (bs, 2H), 3.67 (bs, 2H), 1.60, (bs, 2H), 1.54 (bs, 2H), 1.41 (m, 2H), 1.25 (bs, 2H), 1.13 (m, 2H) ppm.

15 Part B. The product from part A (1.09 g, 3.03 mmol) was dissolved in toluene (10.0 mL) and [2,4-bis(4-methoxyphenyl)-1,3-dithia-2,4-diphosphetane-2,4-disulfide] (0.9 g, 2.2 mmol) was added. The reaction was heated to 90°C for 1.5 h and cooled. An additional 0.5 g (1.23 mmol)
20 of [2,4-bis(4-methoxyphenyl)-1,3-dithia-2,4-diphosphetane-2,4-disulfide] was added and heated for 12 h. The reaction mixture was concentrated and purified via flash chromatography (silica, EtOAc:Hexanes=3:1) to yield [1-(4-Iodo-phenyl)-cyclopropyl]-piperidin-1-yl-methanethione
25 (0.92 g, yield: 88%). LC/MS (ESI⁺) 372.0 (M+H)⁺, t_R=6.56 min (5-98% CH₃CN/H₂O in a 10-min run).

Part C. The product from Part B (0.92 g, 2.4 mmol) was treated with neat methyl iodide (2.00 mL, 32.1 mmol) at rt
30 and allowed to stir under N₂ for 48 h. The reaction was concentrated and stripped (3x) with methanol to provide a yellow solid of 1-{{[1-(4-iodo-phenyl)-cyclopropyl]-methylsulfanyl-methylene}-piperidinium; iodide (0.51 g, yield: 41%).

35

Part D. The product from Part C (0.51 g, 0.99 mmol) was dissolved in methanol (3.0 mL) and ethylenediamine (0.1 mL, 1.49 mmol) was added dropwise at rt. After 2h, reaction mixture was concentrated to dryness and purified via flash chromatography (silica, 100% EtOAc, then 0.5% Et₃N:10% MeOH:CH₂Cl₂) to yield 2-[1-(4-iodo-phenyl)-cyclopropyl]-4,5-dihydro-1H-imidazole (0.20 g, yield: 66%). ¹H NMR (CD₃OD, 300 MHz) δ 7.68 (d, J=8.4 Hz, 2H), 7.14 (d, J=8.5 Hz, 2H), 3.61 (s, 4H), 1.48 m, 2H), 1.25 (s, 2H) ppm.

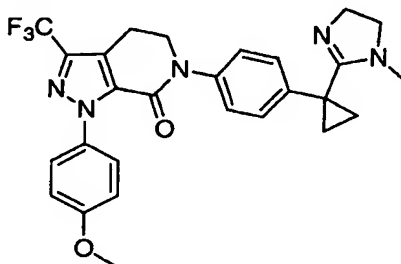
10

Part E. The product from Part D (0.093 g, 0.298 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.096 g, 0.309 mmol) and 2-[1-(4-iodo-phenyl)-cyclopropyl]-4,5-dihydro-1H-imidazole (0.093 g, 0.298 mmol) were coupled by the usual procedure. Purification was accomplished using flash chromatography (silica, 100% EtOAc then 0.5% Et₃N:10% MeOH:CH₂Cl₂) to give product (35 mg, yield: 38%). LC/MS (ESI⁺) 496.6 (M+H)⁺, t_R=2.16 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (CD₃OD, 300 MHz) δ 7.47 (AA'BB', J=8.6 Hz, 4H), 7.30 (d, J=8.4 Hz, 2H), 6.98 (d, J=9.1 Hz, 2H), 4.11 (t, J=6.6 Hz, 2H) 3.82 (s, 3H), 3.52 (s, 4H), 3.15 (t, J=6.6 Hz, 2H), 1.44 (m, 2H), 1.16 (m, 2H) ppm.

25

Example 145

1-(4-Methoxyphenyl)-6-{4-[1-(1-methyl-4,5-dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



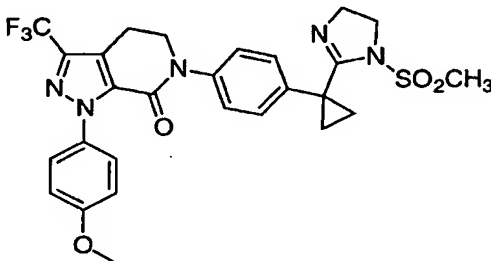
The title compound was obtained following the same sequence as those in Example 145 but using N-methylethylenediamine instead of ethylene diamine. LC/MS (ESI⁺) 510.6 (M+H)⁺, t_R=2.75 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR

(CD₃Cl₃, 300 MHz) δ 7.45 (d, J=8.8 Hz, 2H), 7.27 (m, 4H), 6.92 (d, J=9.2 Hz, 4H), 4.12 (m, 2H), 3.85 (m, 2H), 3.81 (s, 3H), 3.16 (t, J=6.6 Hz, 2H), 2.89 (s, br, 3H), 1.59 (m, 2H), 1.28 (m, 2H) ppm.

10

Example 146

6-(4-[1-(1-Methanesulfonyl-4,5-dihydro-1H-imidazol-2-yl)-cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one

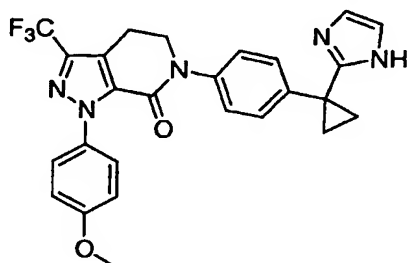


15 The product from Example 144 (0.017 g, 0.034 mmol) from was dissolved in CH₂Cl₂ (0.3 mL). Et₃N (0.01mL, 0.07 mmol) and MsCl (0.07 mL, 0.09 mmol) were added at rt under N₂. The reaction was stirred for 12 h, concentrated to dryness, and purified via flash chromatography (silica, 100% EtOAc then 20 0.5% Et₃N:10% MeOH:CH₂Cl₂) to afford the title compound.

LC/MS (ESI⁺) 574.4 (M+H)⁺, t_R=2.84 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR ((CD₃)₂CO, 300 MHz) δ 7.50 (d, J=9.1 Hz, 2H), 7.33 (m, 4H), 6.98 (d, J=9.1 Hz, 2H), 4.16 (t, J=6.6 Hz, 2H), 3.83 (m, 7H), 3.16 (t, J=6.4 Hz, 2H), 2.42 25 (s, 3H), 1.47 (m, 2H), 1.18 (m, 2H) ppm.

Example 147

6-(4-[1-(1*H*-Imidazol-2-yl)cyclopropyl]phenyl)-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one



5

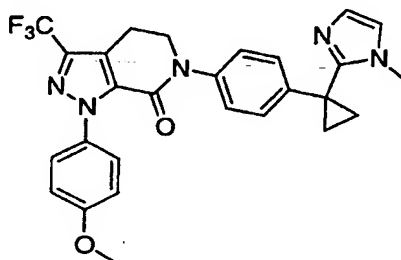
Part A. H-(Boc)-DAP-OMe was dissolved in CH₂Cl₂ (2.0 mL) and TFA (1.0 mL) was added. The reaction was allowed to stir for 2h. The reaction was concentrated and stripped with CHCl₃ (10 mL x 3). The reaction mixture was re-diluted with MeOH (6.0 mL) and K₂CO₃ (spatula tip) added. The product from Part C in Example 142 (0.12 g, 0.234 mmol) was added and the reaction was heated to 65°C for 2h. The reaction was concentrated and purified via flash chromatography (silica, EtOAc-10% MeOH/CH₂Cl₂) to afford 2-[1-(4-Iodo-phenyl)-cyclopropyl]-4,5-dihydro-3*H*-imidazole-4-carboxylic acid methyl ester. LC/MS (ESI⁺) 371.0 (M+H)⁺, t_R=2.17 min (10-90% CH₃CN/H₂O in a 4-min run).

Part B. 1-(4-Methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one (45.0 mg, 0.145 mmol) and product from Part A (36.0 mg, 0.098 mmol) were dissolved in DMSO (0.5 mL). K₂CO₃ (10.0 mg, 0.723 mmol) was added followed by 1,10-phenanthroline (spatula tip) and copper iodide (spatula tip). The reaction was heated to 110°C for 12 h. The reaction was diluted with EtOAc and washed with H₂O (2x) and brine. Organic was dried over NaSO₄, filtered, and concentrated. The reaction was purified via flash chromatography (silica, EtOAc-10% MeOH/1% EtN₃/CH₂Cl₂). Side product obtained from loss of

CO₂Me under the basic conditions to form the title compound. LC/MS (ESI⁺) 494.2 (M+H)⁺, t_R=2.21 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR ((CD₃)₂CO, 300 MHz) δ 7.50 (d, J=9.2 Hz, 2H), 7.32 (m, 4 H), 6.98 (d, J=9.1 Hz, 2H),
 5 6.83 (s, 2H), 4.16 (t, J=6.6 Hz, 2H), 3.83 (s, 3H), 3.16 (t, J=6.2 Hz, 2H), 1.47 (m, 2H), 1.18 (m, 2H) ppm.

Example 148

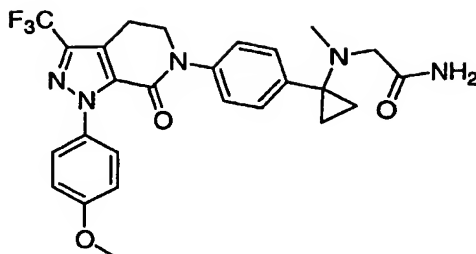
1-(4-Methoxyphenyl)-6-{4-[1-(1-methyl-1H-imidazol-2-yl)-
 10 cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-
 pyrazolo[3,4-c]pyridin-7-one



The product from Example 144 (0.08 g, 0.18 mmol) was dissolved in 1,4-dioxane (2.0 mL) and KMnO₄ (spatula tip)
 15 was added. The reaction was heated to 80°C for 12 h at rt. The reaction was filtered, concentrated, and purified via flash chromatography (silica, 100% EtOAc then 0.5% Et₃N:10% MeOH/CH₂Cl₂ then 100% MeOH) to yield the title compound
 (0.06 g, yield: 72%). ¹H NMR δ 7.44 (d, J=9.1 Hz, 2H), 7.24
 20 (d, J=8.7 Hz, 2H), 6.98 (m, 4H), 4.07 (dd, J=2.9, 3.7 Hz, 2H), 3.81 (s, 3H), 3.48 (s, 2H), 3.12 (t, J=6.2 Hz, 2H), 1.44 (m, 2H), 1.21 (t, J=7.3 Hz, 2H) ppm.

Example 149

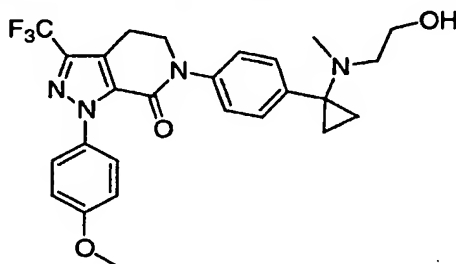
25 2-[(1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-cyclopropyl)-methyl-amino]-acetamide



The product of Example 130 (50 mg, 0.11 mmol) was stirred in DMF (0.3 mL). K_2CO_3 (45 mg, mmol, 0.33 mmol, 3 eq) and chloroacetamide (20 mg, 0.21 mmol, 2 eq) were added. The mixture was stirred at 70°C for 2h. EtOAc was added, washed with H_2O and brine, dried over $MgSO_4$, filtered, and concentrated. The residue was purified by Silica gel purification to yield the title compound. LC/MS (ESI⁺) 514.8 (M+H)⁺, t_R =2.00 min (10-90% CH_3CN/H_2O in a 4-min run). ¹H NMR ($CDCl_3$) δ 7.46 (d, J =9.2 Hz, 2H), 7.27 (m, 4H), 6.91 (d, J =8.8 Hz, 2H), 4.12 (t, J =6.6 Hz, 2H), 3.81 (s, 3H), 3.15 (m, 4H), 2.29 (m, 3H), 0.97 (m, 2H), 0.87 (m, 2H) ppm.

Example 150

6-(4-(1-[(2-Hydroxyethyl)-methylamino]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydropyrazolo[3,4-c]pyridin-7-one



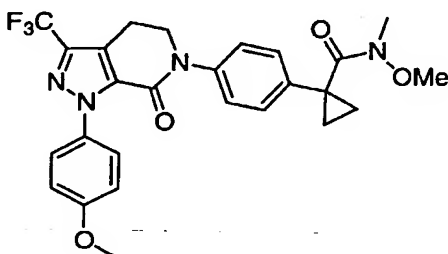
Following a procedure analogous to that used for the preparation of Example 149 but using 2-bromoethanol instead, the title compound was prepared. Silica gel purification yielded the pure desired product. LC/MS (ESI⁺) 501.8 (M+H)⁺, t_R =2.04 min (10-90% CH_3CN/H_2O in a 4-min run). ¹H NMR ($CDCl_3$) δ 7.46 (d, J =9.1 Hz, 2H), 7.29 (m,

4H), 6.92 (d, $J=8.8$, 2H), 4.14 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.51 (m, 2H), 3.16 (t, $J=6.6$ Hz, 2H), 2.64 (m, 2H), 2.23 (s, 3H), 0.95 (s, 3H), 0.83 (m, 2H) ppm.

5

Example 151

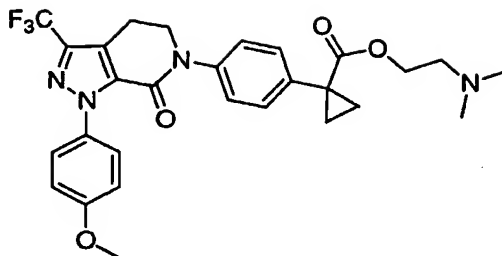
1-(4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropanecarboxylic acid methoxy-methyl-amide



10 1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}cyclopropane carboxylic acid (1.12 g, 2.37 mmol) and N,O-dimethylhydroxylamine hydrochloride (0.255 g, 2.61 mmol) were dissolved in DMF (20.0 mL) and DIEA (2.0 mL, 11.5 mmol) was added dropwise. 1-[3-(Dimethylamino)propyl]-3-ethylcarbodiimide hydrochloride (0.54 g, 2.8 mmol) was added and the reaction was allowed to proceed under nitrogen at rt for 12 h. The reaction was diluted with EtOAc, washed 1N HCl, brine, dried over Na₂SO₄, filtered and
20 concentrated. The reaction was purified via flash chromatography (silica, 100% EtOAc, then 10% MeOH/CH₂Cl₂) to give the title compound. LC/MS (ESI⁺) 515.6 (M+H).

Example 152

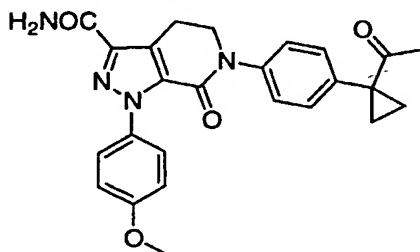
25 **6-[4-(1-Hydroxymethylcyclopropyl)phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide**



The title compound was obtained as a side product in the reaction from Example 151 (0.55 g, 0.96 mmol) was obtained in 40% yield. LC/MS (ESI⁺) 556.4 (M+H)⁺, t_R=1.77 min (10-
 5 90% CH₃CN/H₂O in a 4-min run). ¹H NMR (300 MHz, (CD₃)₂CO) δ
 7.50 (d, J=8.8 Hz, 2H), 7.39 (AA'BB', J=8.8 Hz, 4H), 7.01
 (d, J=8.8 Hz, 2H), 4.19 (t, J=6.6 Hz, 2H), 3.83 (s, 3H),
 3.76 (m, 2H), 3.20 (m, 4H), 2.85 (s, 3H), 2.63 (s, 6H),
 1.52 (m, 2H), 1.31 (m, 2H), 1.09 (t, J=7.0 Hz, 2H), 0.81
 10 (t, J=6 Hz, 1H) ppm.

Example 153

6-[4-(1-Acetyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-
 15 oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
 c]pyridine-3-carboxylic acid amide

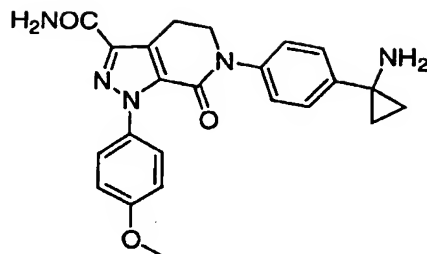


Following a procedure analogous to that used in Example 138, the title compound was prepared. LC/MS (ESI⁺) 445.6 (M+H).

20

Example 154

6-[4-(1-Aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
 4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic
 acid amide, trifluoroacetic acid salt



Part A. [1-(4-Iodo-phenyl)-cyclopropyl]-carbamic acid tert-butyl ester (0.34 g, 0.95 mmol) and 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.30 g, 0.97 mmol) were stirred in DMSO (1 mL). K_2CO_3 (0.25 g, 1.81 mmol), CuI (87 mg, 0.46 mmol) and 1,10-phenanthroline (83 mg, 0.46 mmol) were added. The resulting mixture was heated at 120°C for 2.5 h. After cooling, it was extracted with EtOAc (2x), washed with H_2O and brine, dried over $MgSO_4$, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, CH_2Cl_2 :EtOAc=1:1, then EtOAc) to give 6-[4-(1-tert-butoxycarbonylamino-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.24 g, yield: 46%).

Part B. The product from Part A underwent the same reaction as used in Part E of Example 67 to yield (1-{4-[3-carbamoyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-carbamic acid tert-butyl ester. HRMS $C_{28}H_{32}N_5O_5$ ($M+H$)⁺ 518.2388 calcd for 518.2325. 1H NMR ($CDCl_3$) δ 7.47 (d, $J=9.2$ Hz, 2H), 7.23 (m, 4H), 6.93 (d, $J=9.1$ Hz, 2H), 6.85 (s, br, 1H), 5.52 (s, br, 1H), 5.26 (s, br, 1H), 4.08 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.36 (t, $J=6.6$ Hz, 2H), 1.42 (s, br, 9H), 1.24 (m, 2H), 1.18 (m, 2H) ppm.

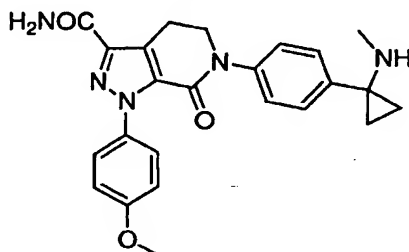
Part C. The product from Part B (54 mg, 0.104 mmol) was stirred in CH_2Cl_2 (2 mL) and TFA (1 mL) at rt for 30 min.

After evaporation, the residue was purified by reverse phase HPLC to afford the title compound (40 mg, yield: 91.7%). HRMS $C_{23}H_{23}N_5O_3$ (M+H)⁺ 418.1908 calcd for 418.1879.

5

Example 155

1-(4-Methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



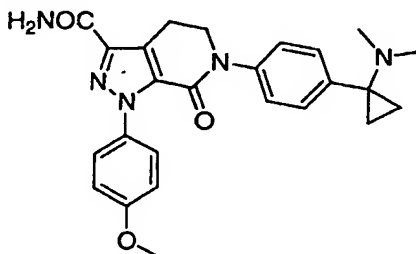
10

Following a procedure analogous to that used in Example 166, the title compound was prepared. LC/MS (ESI⁺) 432.6 (M+H)⁺, t_R =1.74 min (10-90% CH₃CN/H₂O in a 4-min run). ¹H NMR (acetone-*d*₆) δ 7.65 (d, J =8.5 Hz, 2H), 7.49 (m, 4H), 7.33 (s, br, 1H), 6.97 (d, J =9.2 Hz, 2H), 6.74 (s, br, 1H), 5.69 (s, 1H), 4.15 (t, J =6.6 Hz, 2H), 3.82 (s, 3H), 3.28 (t, J =6.6 Hz, 2H), 2.63 (s, 3H), 1.63 (m, 2H), 1.19 (m, 2H) ppm.

20

Example 156

6-[4-(1-Dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt

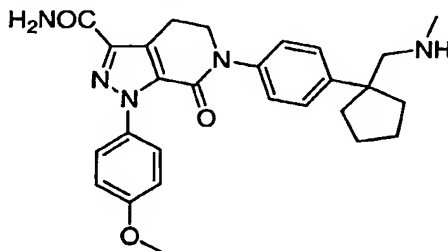


25

The product of Example 154, HOAc (0.05 mL), and aqueous paraformaldehyde (0.3 mL) were stirred in CH₃CN (1.5 mL) at rt for 15 min. NaBH₃CN (60 mg) was added. The mixture was stirred at rt for 2h. H₂O was added. The organic solvent
5 was evaporated. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run). HRMS C₂₅H₂₈N₅O₃ (M+H)⁺ 446.2178 calcd for 446.2193. ¹H NMR (acetone-*d*₆) δ 7.67 (d, *J*=8.4 Hz, 2H), 7.51 (AA'BB', *J*=8.8 Hz, 4H), 6.98 (d, *J*=9.1 Hz, 2H), 4.17 (t, *J*=6.6 Hz, 2H), 3.83 (s, 3H), 3.29 (t, *J*=6.6 Hz, 2H), 2.77 (s, 6H), 1.67 (m, 2H), 1.15 (m, 2H)
10 ppm.

Example 157

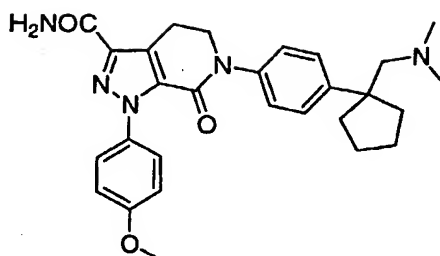
6-[4-(1-Methylaminomethylcyclopentyl)phenyl]-1-(4-methoxy-
15 phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4*c*]pyridine-
3-carboxylic acid amide, trifluoroacetic acid salt



Following an analogous procedures as those used in Examples 27 and 67, the title compound was prepared. HRMS C₂₇H₃₂N₅O₃
20 (M+H)⁺ 474.2533 calcd for 474.2506.

Example 158

6-[4-(1-Dimethylaminomethylcyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-
25 c]pyridine-3-carboxylic acid amide, trifluoroacetic acid
salt

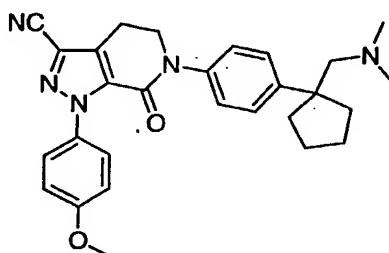


Following an analogous procedures as those used in Examples 28 and 68, the title compound was prepared. LC/MS (ESI⁺) 488.6 (M+H)⁺, t_R =1.77 min (10-90% CH₃CN/H₂O in a 4-min run).

5

Example 159

6-[4-(1-Dimethylaminomethylcyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



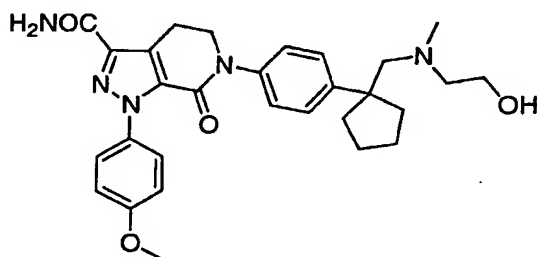
10

Following an analogous procedures as those used in Examples 28 and 75, the title compound was prepared. HRMS C₂₈H₃₂N₅O₃ (M+H)⁺ 470.2577 calcd for 470.2557.

15

Example 160

6-[4-(1-[(2-Hydroxyethyl)methylaminomethyl]cyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



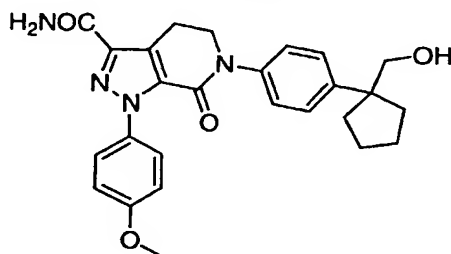
20

Following a procedure analogous to that used in Example 150 but using the product of Example 157 as the starting material, the title compound was prepared. The product was purified by RP-prep LC-MS (5-98% CH₃CN/H₂O in a 10-min run).

5 LC/MS (ESI⁺) 518.8 (M+H). ¹H NMR (acetone-d₆) δ 7.56 (d, J=8.8 Hz, 2H), 7.51 (d, J=9.1 Hz, 2H), 7.39 (d, J=8.8 Hz, 2H), 6.97 (d, J=9.2 Hz, 2H), 4.12 (t, J=6.6 Hz, 2H), 3.82 (s, 3H), 3.78 (m, 2H), 3.69 (m, 2H), 3.27 (t, J=6.6 Hz, 2H), 3.15 (t, J=5.0 Hz, 2H), 2.60 (s, 3H), 2.17 (m, 4H),
10 1.81 (m, 2H), 1.64 (m, 2H) ppm.

Example 161

6-[4-(1-Hydroxymethyl-cyclopentyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
15 c]pyridine-3-carboxylic acid amide

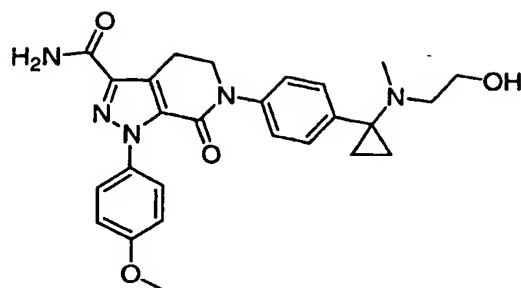


Following a procedure analogous to that used in Example 95, the title compound was prepared. LC/MS (ESI⁺) 461.4.

20

Example 162

6-(4-(1-[(2-Hydroxyethyl)methylamino]cyclopropyl)phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amide, trifluoroacetic acid
salt

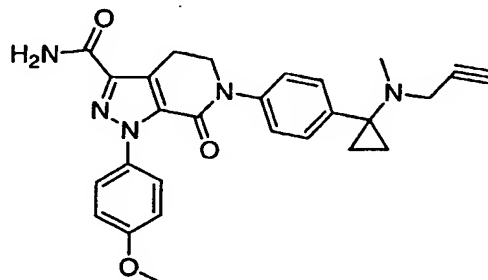


Following a procedure analogous to that used in Example 150 but using the product of Example 155 as the starting material, the title compound was prepared. HRMS $C_{26}H_{30}N_5O_4$

5 (M+H)⁺ 476.2299 calcd for 476.2319.

Example 163

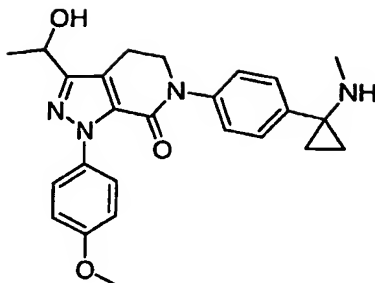
1- (4-Methoxyphenyl) -6- {4- [1- (methyl-prop-2-ynylamino) -
cyclopropyl]phenyl} -7-oxo-4,5,6,7-tetrahydro-1H-
10 pyrazolo[3,4-c]pyridine-3-carboxylic acid amide,
trifluoroacetic acid salt



Following a procedure analogous to that used in Example 162 but using 3-bromo-propyne as the starting material instead
15 of 2-bromoethanol, the title compound was prepared. HRMS
 $C_{27}H_{28}N_5O_3$ (M+H)⁺ 470.2178 calcd for 470.2193.

Example 164

3- (1-Hydroxyethyl) -1- (4-methoxyphenyl) -6- [4- (1-methylamino-
20 cyclopropyl)phenyl] -1,4,5,6-tetrahydro-pyrazolo[3,4-
c]pyridin-7-one, trifluoroacetic acid salt



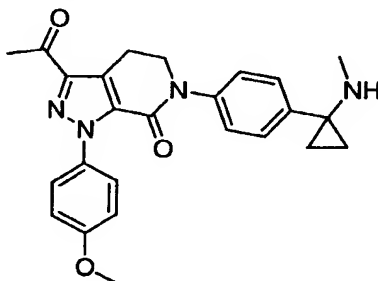
Part A. 6-(4-[1-(tert-Butoxycarbonyl-methyl-amino)-cyclopropyl]-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid
 5 ethyl ester (0.59 g, 1.79 mmol) was stirred in EtOH (15 mL) and 1N NaOH (3 mL) at rt for 1h. After evaporation, the mixture was acidified with citric acid. The mixture was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness in vacuo.
 10 This acid underwent a series of reactions similar to those used in Part E of Example 1 to afford (1-{4-[3-formyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-methyl-carbamic acid tert-butyl ester (yield: 99%). LC/MS (ESI⁺) 461.6 (M+H-t-Bu)⁺, t_R=1.2.97 min (10-90% CH₃CN/H₂O in a 4-min run).
 15

Part B. The product from Part A (0.38 g, 0.74 mmol) was stirred in CH₂Cl₂ (6 mL) at -78°C under N₂. ZnMe₂ (2M in toluene, 0.74 mL, 1.48 mmol) was added dropwise followed by
 20 the addition of TiCl₄ (0.16 mL, 1.07 mmol) dropwise. The reaction was stirred at -78°C for 2h. Saturated NH₄Cl was added. The mixture was extracted with EtOAc, washed with H₂O and brine, dried over MgSO₄, filtered and concentrated. The residue was purified by FCC (silica gel, CH₂Cl₂, then
 25 20% EtOAc in CH₂Cl₂) to yield (1-{4-[3-(1-hydroxy-ethyl)-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-methyl-carbamic acid tert-butyl ester (81 mg, yield: 25%). HRMS C₂₆H₃₀N₅O₄ (M+H)⁺ 533.2778 calcd for 533.2765.

Part C. The product from Part B (10 mg) was stirred in CH_2Cl_2 (1 mL) and TFA (1 mL) at rt for 30 min. After evaporation, the residue was purified by reverse phase HPLC (0-100% CH_3CN in H_2O) to afford the title compound. HRMS $\text{C}_{26}\text{H}_{30}\text{N}_5\text{O}_4$ ($\text{M}+\text{H}$)⁺ 433.2247 calcd for 433.2240. ^1H NMR (CD_3OD) δ 7.59 (d, $J=8.4$ Hz, 2H), 7.44 (d, $J=8.4$ Hz, 2H), 7.38 (d, $J=8.4$ Hz, 2H), 6.94 (d, $J=8.4$ Hz, 2H), 5.00 (q, $J=6.6$ Hz, 1H), 4.10 (m, 2H), 3.83 (s, 3H), 3.15 (t, $J=6.6$ Hz, 2H), 2.58 (s, 3H), 1.56 (d, $J=8.4$ Hz, 4H), 1.39 (m, 2H), 1.28 (m, 2H) ppm. ^{19}F NMR (CD_3OD) δ -77.49 ppm.

Example 165

3-Acetyl-1-(4-methoxyphenyl)-6-[4-(1-methylamino-cyclopropyl)phenyl]-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

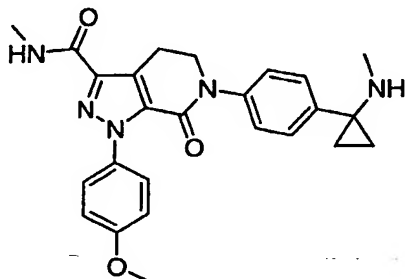


The product from Part B in Example 164 (50 mg, 0.094 mmol), PCC (40 mg, 0.19 mmol), NaOAc (23 mg, 0.28 mmol), and 4Å MS (50 mg) were stirred in CH_2Cl_2 (1 mL) for 4h. The mixture was filtered through Celite®, washed with H_2O (2x), dried over MgSO_4 , filtered and concentrated to dryness. The residue was dissolved in CH_2Cl_2 (4 mL) and TFA (2 mL) at rt for 30 min. After evaporation, the residue was purified by reverse phase HPLC (0-100% CH_3CN in H_2O) to afford the title compound. ^1H NMR (CD_3OD) δ 7.59 (d, $J=8.1$ Hz, 2H), 7.46 (m, 4H), 6.98 (d, $J=9.1$ Hz, 2H), 4.11 (t, $J=6.6$ Hz, 2H), 3.82 (m, 5H), 3.29 (m, 2H), 2.59 (m, 3H), 1.39 (m, 2H), 1.28 (m,

2H) ppm. ^{19}F NMR (CD_3OD) δ -77.51 ppm. HRMS $\text{C}_{25}\text{H}_{27}\text{N}_4\text{O}_3$
($\text{M}+\text{H}$) $^+$ 431.2102 calcd for 431.2084.

Example 166

5 1-(4-Methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid methylamide, trifluoroacetic acid salt

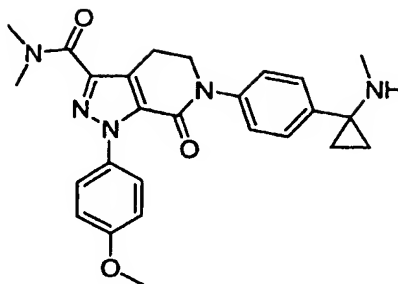


The product from Part A in Example 164 (0.21 g, 0.44 mmol)
10 was stirred in CH_2Cl_2 (5 mL) at 0°C . $(\text{COCl})_2$ (0.1 mL) was
added followed by the addition of 1 drop of DMF. The
mixture was stirred at 0°C for 40 min. The solvents were
evaporated in vacuo. Half of the residue was dissolved in
 CH_2Cl_2 (1 mL) and MeNH_2 (2 M in THF, 0.5 mL) was added. The
15 mixture was stirred at rt for 4h. The solvents were
evaporated. The residue was dissolved in CH_2Cl_2 (15 mL) and
TFA (2 mL). The mixture was stirred at rt for 1h. The
solvents were evaporated. The residue was purified by
reverse phase HPLC (0-100% CH_3CN in H_2O with 0.5% TFA) and
20 lyophilized to dryness. HRMS $\text{C}_{25}\text{H}_{28}\text{N}_5\text{O}_3$ ($\text{M}+\text{H}$) $^+$ 446.2177
calcd for 446.2193. ^1H NMR (CD_3OD) δ 7.59 (d, $J=9.1$ Hz,
2H), 7.47 (m, 4H), 6.96 (d, $J=9.1$ Hz, 2H), 4.17 (t, $J=6.6$
Hz, 2H), 3.82 (m, 5H), 3.29 (m, 2H), 3.17 (m, 2H), 2.92 (m,
3H), 1.61 (m, 2H), 1.34 (m, 2H) ppm.

25

Example 167

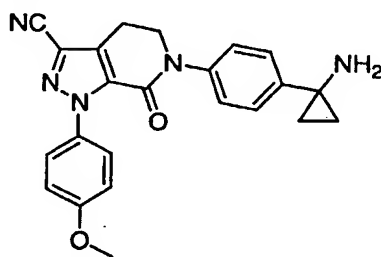
1-(4-Methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid dimethylamide, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 166 but using dimethylamine as the starting material, the title compound was prepared. The product was purified by reverse
 5 phase HPLC (0-100% CH₃CN in H₂O with 0.5% TFA) and lyophilized to dryness. HRMS C₂₆H₃₀O₃N₅ (M+H)⁺ 460.2319 calcd for 460.2349. ¹H NMR (CD₃OD) δ 7.59 (m, 2H), 7.45 (m, 4H), 6.96 (m, 2H), 4.11 (t, J=6.6 Hz, 2H), 3.37 (s, 3H), 3.17 (t, J=6.6 Hz, 2H), 3.12 (s, 3H), 2.59 (s, 3H), 1.39
 10 (m, 2H), 1.29 (m, 2H) ppm. ¹⁹F NMR (CD₃OD) δ -77.56 ppm.

Example 168

6-[4-(1-Aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
 4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
 15 carbonitrile, trifluoroacetic acid salt

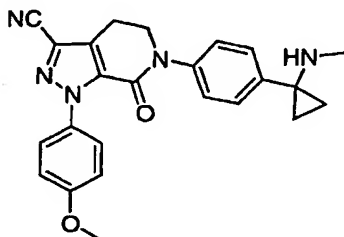


Following a procedure analogous to that used in Example 74 but using the product of Example 154 as the starting material, the title compound was prepared. The product was
 20 purified by reverse phase HPLC (0-100% CH₃CN in H₂O with 0.5% TFA) and lyophilized to dryness. LC/MS(ESI⁺) 400.4 (M+H)⁺. ¹H NMR (acetone-d₆) δ 7.52 (d, J=9.1 Hz, 2H), 7.41 (AA'BB', J=8.0 Hz, 4H), 6.98 (d, J=9.1 Hz, 2H), 4.19 (t,

$J=6.6$ Hz, 2H), 3.83 (s, 3H), 3.20 (t, $J=6.6$ Hz, 2H), 1.69 (m, 2H), 1.51 (m, 2H) ppm.

Example 169

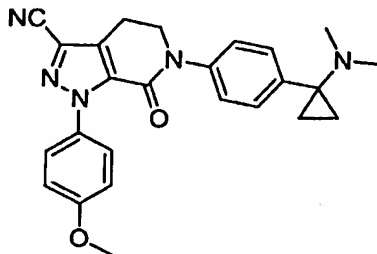
5 1-(4-Methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 74
 10 but using the product of Example 155 as the starting material, the title compound was prepared. HRMS $C_{24}H_{24}O_2N_5$ (M+H)⁺ 414.1900 calcd for 414.1931. ¹H NMR (acetone- d_6) δ 7.68 (d, $J=8.4$ Hz, 2H), 7.49 (AA'BB', $J=9.2$ Hz, 4H), 6.98 (d, $J=9.1$ Hz, 2H), 4.23 (t, $J=6.6$ Hz, 2H), 3.83 (s, 3H),
 15 3.19 (t, $J=6.2$ Hz, 2H), 2.62 (s, 3H), 1.64 (t, $J=6.6$ Hz, 2H), 1.19 (t, $J=6.5$ Hz, 2H) ppm.

Example 170

20 6-[4-(1-Dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt

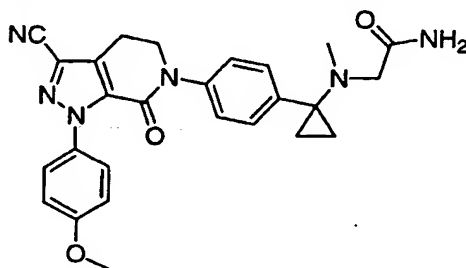


Following a procedure analogous to that used in Example 67
 but using the product of Example 156 as the starting
 25 material, the title compound was prepared. HRMS $C_{25}H_{26}O_2N_5$

(M+H)⁺ 428.2104 calcd for 428.2087. ¹H NMR (acetone-d₆) δ
 7.70 (d, J=8.4 Hz, 2H), 7.55 (AA'BB', J=9.2 Hz, 4H), 7.00
 (d, J=9.1 Hz, 2H), 4.26 (t, J=6.6 Hz, 2H), 3.83 (s, 3H),
 3.22 (t, J=6.2 Hz, 2H), 2.79 (s, 6H), 1.70 (t, J=6.1 Hz,
 5 2H), 1.17 (t, J=6.1 Hz, 2H) ppm.

Example 171

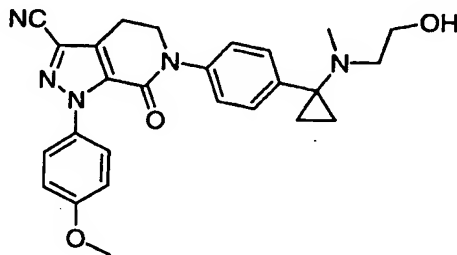
2-[(1-{4-[3-Cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
 tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropyl)-
 10 methylamino]acetamide, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 162
 but using the product of Example 169 as the starting
 material, the title compound was prepared. The product was
 15 purified by reverse phase HPLC (0-100% CH₃CN in H₂O with
 0.5% TFA) and lyophilized to dryness. LC/MS (ESI) 471.6
 (M+H), t_R=2.14 min (10-90% CH₃CN in H₂O in a 4-min run).

Example 172

20 6-(4-{1-[(2-Hydroxyethyl)methylamino]cyclopropyl}phenyl)-1-
 (4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
 c]pyridine-3-carbonitrile, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 162
 25 but using the product of Example 169 and 2-bromoethanol as

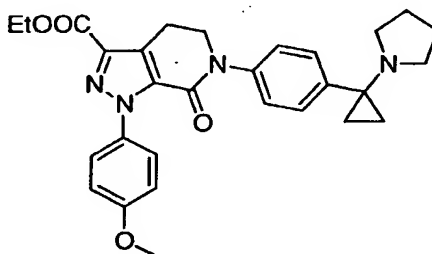
the starting materials, the title compound was prepared. The product was purified by reverse phase HPLC (0-100% CH₃CN in H₂O with 0.5% TFA) and lyophilized to dryness.

HRMS C₂₆H₂₈N₅O₃ (M+H)⁺ 458.2196 calcd for 458.2193. ¹H NMR

(CD₃OD) δ 7.68 (m, 2H), 7.46 (m, 4H), 6.98 (m, 2H), 4.17 (t, J=6.6 Hz, 2H), 3.82 (m, 5H), 3.29 (m, 2H), 3.17 (m, 2H), 2.92 (m, 3H), 1.61 (m, 2H), 1.34 (m, 2H) ppm. ¹⁹F NMR (CD₃OD) δ -77.56 ppm.

Example 173

1-(4-Methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester, trifluoroacetic acid salt

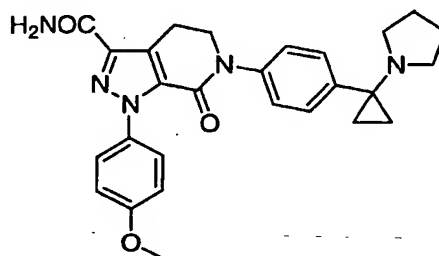


6-[4-(1-Amino-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (95 mg, 0.21 mmol), 1,3-dibromo-propane (0.03 mL, excess), and K₂CO₃ (100 mg, excess) were heated in DMF at 80°C for 24 h. EtOAc was added. The mixture was washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated. The residue was purified by FCC (Silica gel, EtOAc, then 15% MeOH in EtOAc) to yield the title compound (89 mg, yield: 83.5%). HRMS C₂₉H₃₃O₄N₄ (M+H)⁺ 501.2489 calcd for 501.2503. ¹H NMR (CDCl₃) δ 7.47 (d, J=8.8 Hz, 2H), 7.27 (AA'BB', J=8.8 Hz, 4H), 6.90 (d, J=8.9 Hz, 2H), 4.45 (q, J=7.1 Hz, 2H), 4.12 (t, J=6.6 Hz, 2H), 3.80 (s, 3H), 3.31 (t, J=6.6 Hz, 2H),

2.52 (m, 4H), 1.61 (m, 4H), 1.43 (t, $J=7.1$ Hz, 2H), 0.97 (m, 2H), 0.77 (m, 2H) ppm.

Example 174

5 **1-(4-Methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt**

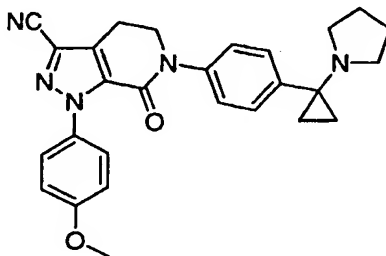


10 Following a procedure analogous to that used in Example 67 but using the product of Example 173 as the starting material, the title compound was prepared. The product was purified by flash chromatography (silica gel, EtOAc, then 15% MeOH in EtOAc). LC/MS(ESI⁺) 472.6 (M+H)⁺, $t_R=2.02$ min
15 (10-90% CH₃CN in H₂O in a 4-min run). ¹H NMR (CDCl₃) δ 7.39 (d, $J=8.8$ Hz, 2H), 7.21 (AA'BB', $J=8.8$ Hz), 6.86 (d, $J=9.1$ Hz, 2H), 4.04 (t, $J=6.9$ Hz, 2H), 3.74 (s, 3H), 3.30 (t, $J=6.6$ Hz, 2H), 2.47 (m, 4H), 1.54 (m, 4H), 0.93 (m, 2H), 0.71 (m, 2H) ppm.

20

Example 175

1-(4-Methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt

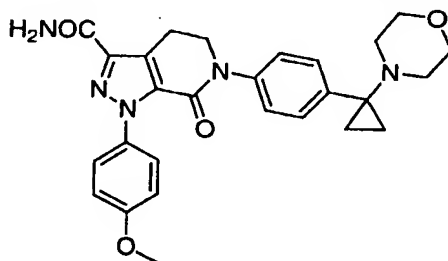


25

Following a procedure analogous to that used in Example 74 but using the product of Example 174 as the starting material, the title compound was prepared. The product was purified by flash chromatography (silica gel, EtOAc, then 5 15% MeOH in EtOAc). LC/MS(ESI⁺) 454.6 (M+H)⁺, t_R=2.27 min (10-90% CH₃CN in H₂O in a 4-min run). ¹H NMR (CDCl₃) δ 7.39 (d, J=8.8 Hz, 2H), 7.28 (d, J=8.4 Hz, 2H), 7.20 (d, J=8.5 Hz), 6.86 (d, J=9.2 Hz, 2H), 4.09 (t, J=6.6 Hz, 2H), 3.75 (s, 3H), 3.10 (t, J=6.6 Hz, 2H), 2.63 (m, 4H), 1.62 (m, 10 4H), 1.17 (m, 2H), 0.78 (m, 2H) ppm.

Example 176

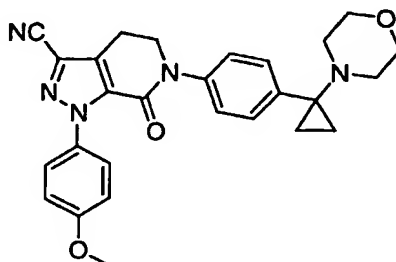
15 1-(4-Methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



The product of Example 153 (45 mg, 0.108 mmol) and 1-bromo-2-(2-bromo-ethoxy)-ethane (0.25 mL), Et₃N (0.25 mL) were 20 heated in DMF (1 mL) at 65°C for 3h under N₂. The mixture was evaporated, and the residue was purified by reverse phase HPLC (0-100% CH₃CN in H₂O with 0.5% TFA), and lyophilized to dryness. LC/MS(ESI⁺) 488.6 (M+H)⁺, t_R=1.97 min (10-90% CH₃CN in H₂O in a 4-min run). ¹H NMR (acetone- 25 d₆) δ 7.51 (d, J=9.2 Hz, 2H), 7.49 (m, 4H), 6.97 (d, J=8.8 Hz, 2H), 4.12 (t, J=6.6 Hz, 2H), 3.83 (s, 3H), 3.27 (m, 4H), 3.29 (t, J=6.6 Hz, 2H), 2.63 (m, 4H), 1.08 (m, 2H), 0.83 (m, 2H) ppm.

Example 177

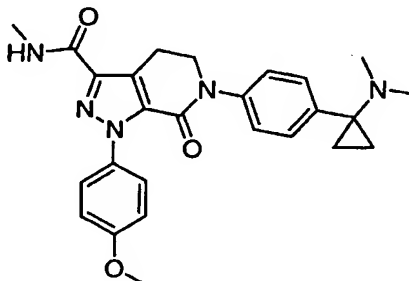
1-(4-Methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 175 but using the product of Example 168 as the starting material, the title compound was prepared. The product was purified by flash chromatography (silica gel, EtOAc, then 10% MeOH in EtOAc). LC/MS(ESI⁺) 470.6 (M+H)⁺. ¹H NMR (CDCl₃) δ 7.46 (d, J=9.2 Hz, 2H), 7.26 (m, 4H), 6.93 (d, J=9.2 Hz, 2H), 4.16 (t, J=6.6 Hz, 2H), 3.82 (s, 3H), 3.75 (m, 2H), 3.65 (m, 4H), 3.50 (t, J=6.3, 2H), 3.18 (t, J=6.6 Hz, 2H), 0.94 (m, 2H), 0.79 (m, 2H) ppm.

Example 178

6-[4-(1-Dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid methylamide, trifluoroacetic acid salt

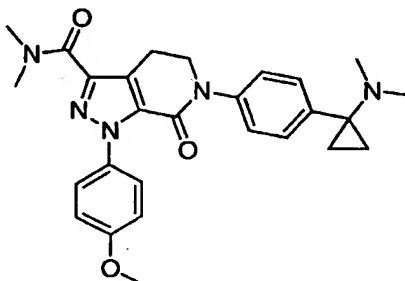


The product was purified by flash chromatography (silica gel, EtOAc, then 10% MeOH in EtOAc). HRMS C₂₆H₃₀N₅O₃ (M+H)⁺

460.2319 calcd for 460.2349. ¹H NMR (acetone-*d*₆) δ 7.49 (d, *J*=9.1 Hz, 2H), 7.30 (m, 4H), 6.95 (d, *J*=8.8 Hz, 2H), 4.11 (t, *J*=6.6 Hz, 2H), 3.82 (s, 3H), 3.28 (t, *J*=6.6 Hz, 2H), 2.90 (d, *J*=4.7 Hz 3H), 2.15 (s, 6H), 0.83 (m, 2H), 0.73 (m, 2H) ppm.

Example 179

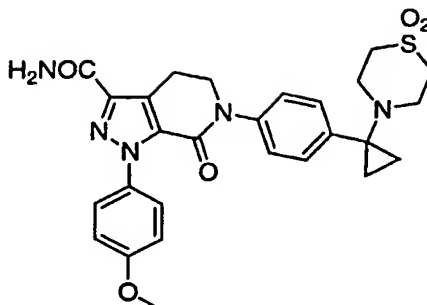
6-[4-(1-Dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxylic acid dimethylamide, trifluoroacetic acid salt



Following a procedure analogous to that used in Example 166, the title compound was prepared. The product was purified by flash chromatography (silica gel, EtOAc, then 10% MeOH in EtOAc). LC/MS(ESI⁺) 474.6 (M+H)⁺, *t*_R=6.08 min (10-90% CH₃CN in H₂O in a 4-min run). ¹H NMR (acetone-*d*₆) δ 7.50 (d, *J*=8.8 Hz, 2H), 7.32 (m, 4H), 6.95 (d, *J*=9.1 Hz, 2H), 4.10 (t, *J*=6.6 Hz, 2H), 3.82 (s, 3H), 3.38 (s, 3H), 3.18 (t, *J*=6.6 Hz, 2H), 3.04 (s, 3H), 2.19 (s, 6H), 0.87 (m, 2H), 0.75 (m, 2H) ppm.

Example 180

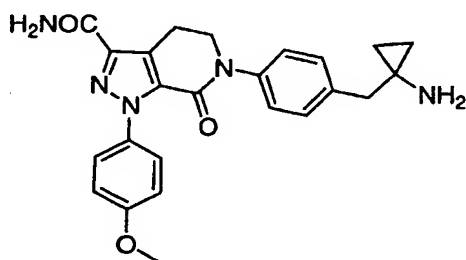
6-[4-[1-(1,1-Dioxo-1λ⁶-thiomorpholin-4-yl)cyclopropyl]phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



The product from Example 153 (90 mg, 0.22 mmol) was stirred in MeOH (1 mL) in a Pyrex® tube. Vinyl sulfone (0.1 mL) was added followed by the addition of Et₃N (0.2 mL). The tube was capped. The mixture was stirred at rt for 1h, and heated at 40-50°C for 1.5 h. After cooling, the solvents were evaporated. The residue was purified by reverse phase HPLC (0-100% CH₃CN in H₂O) and lyophilized to afford the desired product. LC/MS ESI 536.4 (M+H), *t_R*=2.49 (10-90% CH₃CN in H₂O in a 4-min run).

Example 181

6-[4-(1-Aminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



Part A. 4-Iodophenylacetonitrile (0.90 g, 3.70 mmol) was stirred in Et₂O (7 mL) at rt under N₂. Ti(O-*i*Pr)₄ (1.20 mL, 1.1 eq) was added followed by dropwise addition of EtMgBr (2.0 M in Et₂O, 2.5 mL, 2.0 eq) at rt. The reaction mixture was stirred at rt for 0.5 h. BF₃·Et₂O (0.94 mL, 2.0 eq) was added dropwise within 2 mins. The mixture was stirred at rt for 20 min. LC/MS showed one peak corresponding to the desired product. 1N NaOH (ca. 3 mL)

was added. It was extracted with Et₂O (2 x), washed with H₂O, dried over MgSO₄, filtered, and concentrated to dryness to give 1-(4-iodo-benzyl)-cyclopropylamine (0.60 g, yield: 60%). LC/MS (ESI⁺) 274.2 (M+H), t_R=1.61 min (10-90% CH₃CN in H₂O in a 4-min run).

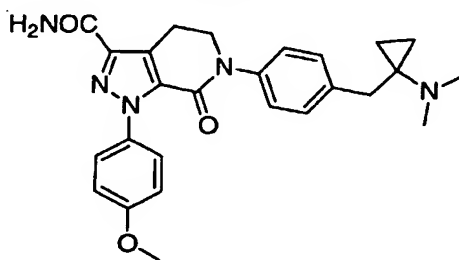
Part B. The product from Part A (0.60 g, 2.2 mmol) was stirred in CH₂Cl₂ (8 mL) at rt under N₂. (Boc)₂O (0.57 g, 1.2 eq) was added followed by the addition of DIEA (0.61 mL, 1.5 eq). The mixture was stirred at rt for 3h. H₂O was added, the mixture was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, and concentrated to dryness to yield [1-(4-iodo-benzyl)-cyclopropyl]-carbamic acid tert-butyl ester (0.61 g, 75%). LC/MS (ESI⁺) 318.0 (M-(t-Bu)+H), t_R=2.84 min (10-90% CH₃CN in H₂O in a 4-min run).

Part C. The product of Part B (0.18 g, 0.48 mmol) and 1-(4-methoxy-phenyl)-7-oxo-3a,4,5,6,7,7a-hexahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.15 g, 0.47 mmol) were stirred in DMSO (1.5 mL) under N₂. K₂CO₃ (0.13 g, 1.0 mmol, 2.1 eq) was added, followed by the addition of CuI (0.050 g, 0.26 mmol) and 1,10-phenanthroline (0.048 g, 0.26 mmol). The mixture was heated at 120°C for 3h. After cooling, it was extracted with EtOAc (2x), washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, CH₂Cl₂:EtOAc=1:1, then EtOAc) to give 6-[4-(1-tert-butoxycarbonylamino-cyclopropylmethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-3a,4,5,6,7,7a-hexahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.11 g, yield: 45%). LC/MS (ESI⁺) 505.2 (M+H-t-Bu), t_R=2.75 min (35-98% CH₃CN in H₂O in a 6-min run).

Part D. The product from Part C (90 mg, 0.16 mmol) was stirred in saturated NH_3 in ethylene glycol at 80°C in a Pyrex® tube for 4h. The cooled mixture was diluted with
5 H_2O , and extracted with EtOAc (2x). The organics were rinsed with H_2O and brine, dried over MgSO_4 , filtered, and concentrated to dryness. The residue was dissolved in CH_2Cl_2 (5 mL), and TFA (3 mL) was added. The mixture was stirred at rt for 20 min. The solvent was evaporated, and
10 the residue was purified by reverse phase HPLC (0-100% CH_3CN in H_2O) to afford pure title compound (45 mg, yield: 65.2%). ^1H NMR (CDCl_3) δ 7.47 (d, $J=8.8$ Hz, 2H), 7.21 (AA'BB', $J=8.4$ Hz, 4H), 6.93 (d, $J=9.2$ Hz, 2H), 4.10 (t, $J=6.6$ Hz, 2H), 3.82 (s, 3H), 3.37 (t, $J=6.6$ Hz, 2H), 2.82
15 (s, br, 2H), 0.75 (s, 4H) ppm. LC/MS (ESI⁺) 432.6 (M+H), $t_R=0.36$ min (35-98% CH_3CN in H_2O in a 6-min run).

Example 182

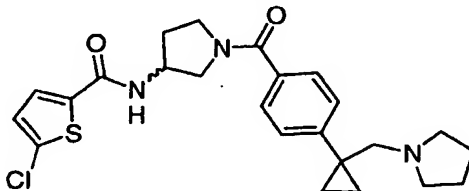
20 6-[4-(1-Dimethylaminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide, trifluoroacetic acid salt



Following the same procedure as shown in Example 155, the
25 title compound was prepared. LC/MS (ESI⁺) 460.6 (M+H)⁺, $t_R=2.14$ min (10-90% CH_3CN in H_2O in a 4-min run).

Example 183

5-Chloro-thiophene-2-carboxylic acid {1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide, trifluoroacetic acid salt



5
Part A. 1-(4-Chlorocarbonyl-phenyl)-cyclopropanecarboxylic acid methyl ester (0.78 g, 3.28 mmol), pyrrolidin-3-yl-carbamic acid tert-butyl ester (0.60 g, 3.22 mmol) and DIEA (1.18 mL, 6.44 mmol) were stirred in CH₂Cl₂ (10 mL) at rt
10 under N₂ overnight. H₂O was added. The mixture was extracted with EtOAc, washed with brine, dried over MgSO₄, filtered, dried in vacuo. The residue was dissolved in MeOH (20 mL) and 1N NaOH (10 mL) was added. The reaction was heated at 50°C for 2.5h. The solvents were evaporated.
15 The residue was extracted with Et₂O, the H₂O layer was acidified with citric acid, and extracted with Et₂O (2x), washed with brine, dried over MgSO₄, filtered, and concentrated to dryness to yield 1-[4-(3-tert-butoxycarbonylamino-pyrrolidine-1-carbonyl)-phenyl]-cyclopropanecarboxylic acid methyl ester (1.08 g, yield:
20 83.0%).

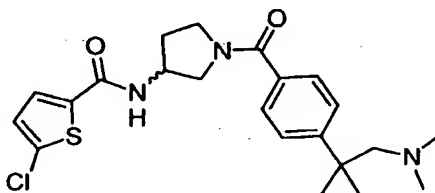
Part B. The product from Part A was treated with an analogous sequence as used in Part E and Part F of Example
25 1 but using pyrrolidine as the starting material and {1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-carbamic acid tert-butyl ester was obtained.

Part C. The product from Part B was stirred in CH₂Cl₂ (20
30 mL) and TFA (10 mL) at rt for 1h. The solvents were evaporated to dryness. Part of the amine (ca. 20 mg) was

dissolved in DMF (0.5 mL). 5-Chloro-thiophenecarbocyclic acid (10 mg) was added followed by the addition of HATU (30 mg) and DIEA (0.03 mL). The reaction was stirred at rt overnight. It was purified via preparative LC/MS (5-98% CH₃CN in H₂O) to afford the desired title compound (12.1 mg, yield: 43.5 %). LC/MS(ESI⁺) 458.6 (M+H), t_R=2.42 min (10-90% CH₃CN in H₂O in a 4-min run).

Example 184

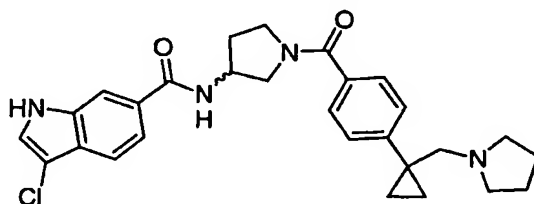
10 **5-Chloro-thiophene-2-carboxylic acid (1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl)-amide, trifluoroacetic acid salt**



Following a procedure analogous to that used in Example 183, the title compound was prepared. LC/MS(ESI⁺) 405.2 (M+H), t_R=2.61 min (10-90% CH₃CN in H₂O in a 4-min run).

Example 185

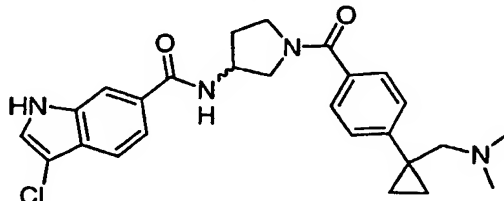
20 **3-Chloro-1H-indole-6-carboxylic acid (1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl)-amide, trifluoroacetic acid salt**



Following a procedure analogous to that used in Example 183, the title compound was prepared. LC/MS (ESI⁺) 491.4 (M+H).

Example 186

3-Chloro-1H-indole-6-carboxylic acid (1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl)-amide, trifluoroacetic acid salt



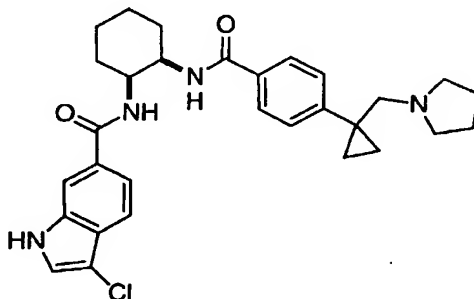
5

Following a procedure analogous to that used in Example 183, the title compound was prepared. LC/MS (ESI⁺) 465.4 (M+H).

10

Example 187

3-Chloro-1H-indole-6-carboxylic acid (2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl)-amide, trifluoroacetic acid salt



15 Part A. 1-(4-Chlorocarbonyl-phenyl)-cyclopropanecarboxylic acid methyl ester (0.66 g, 2.77 mmol) was stirred in CH₂Cl₂ (10 mL) at rt under N₂. 1,2-Cis-diamino-cyclohexane (0.66 mL, 2.0 eq) was added as one portion. The mixture was stirred for 10 min. Diluted HCl was added. The mixture
20 was extracted with EtOAc (2x). The aqueous layer was basified with conc. NaOH, extracted with EtOAc (2x). The organic layers were washed with H₂O and brine, dried over MgSO₄, filtered, and concentrated to afford 1-[4-(2-amino-cyclohexylcarbamoyl)-phenyl]-cyclopropanecarboxylic acid

methyl ester (0.35 g, yield: 39.8%). LC/MS (ESI⁺) 317.4 (M+H).

Part B. The product from Part A (0.15 g, 0.47 mmol) was stirred in DMF (1 mL) at rt. 3-Chloro-1*H*-indole-6-carboxylic acid (0.28 g, 1.36 mmol, 2.9 eq) and HATU (0.36 g, 0.95 mmol, 2.0 eq) were added followed by the addition of DIEA (0.30 mL, 1.71 mmol, 3.6 eq). The mixture was stirred at rt overnight. H₂O was added. The mixture was extracted with EtOAc (2x). The organic layers were washed with brine, dried over MgSO₄, and concentrated to dryness to afford 1-(4-(2-[(3-chloro-1*H*-indole-6-carbonyl)-amino]-cyclohexylcarbamoyl)-phenyl)-cyclopropanecarboxylic acid methyl ester (0.20 g, yield: 85.7%). LC/MS(ESI⁺) 494.6 (M+H), *t*_R=3.22 min (35-95% CH₃CN in H₂O in a 6-min run).

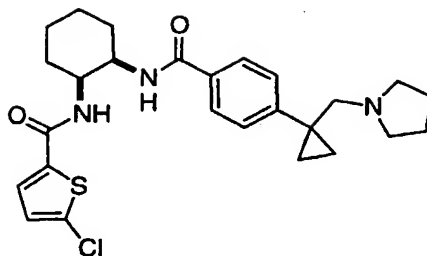
Part C. The product from Part B was subjected to an analogous sequence as used in Part E and Part F of Example 1 but using pyrrolidine as the starting material to afford 3-chloro-6-(2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexylcarbamoyl)-indole-1-carboxylic acid ethyl ester. LC/MS(ESI⁺) 591.6 (M+H).

Part D. The product from Part C was suspended in 4N HCl (20 mL) and heated at 50°C for 1h. The solvent was evaporated. The residue was purified by reverse phase HPLC (0-100% CH₃CN in H₂O) to afford the title compound (32 mg, yield: 33% for Part C and Part D). LC/MS(ESI⁺) 519.4 (M+H), *t*_R=1.85 min (10-90% CH₃CN in H₂O in a 6-min run).

30

Example 188

5-Chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide, trifluoroacetic acid salt

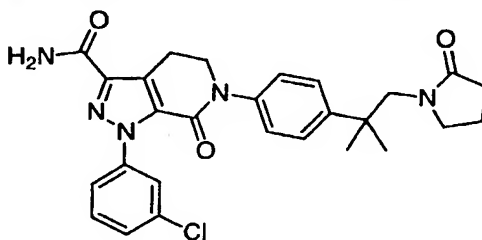


Following a procedure analogous to that used in Example 187, the title compound was prepared. LC/MS (ESI⁺) 466.4 (M+H).

5

Example 189

1-(3-Chloro-phenyl)-6-{4-[1,1-dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



10

Part A. To crude 2-(4-iodo-phenyl)-2-methyl-propionitrile (see Example 96) (1.5 g, 5.5 mmol) in THF (25 mL) at 0°C was added 1M Borane in THF (6 mL, 6 mmol) and the reaction was stirred 2h at rt. The reaction was quenched with water, extracted with ethyl acetate, washed with brine, and dried (Na₂SO₄). The crude residue was treated with 1N HCl and extracted with diethyl ether. The aqueous layer was basified and extracted with ethyl acetate and dried to afford 0.38 g (25%) of a light brown oil. ¹H NMR (CDCl₃) δ 7.59 (d, J=8.4 Hz, 2H), 7.04 (d, J=8.4 Hz, 2H), 2.70 (s, 2H), 1.21 (s, 6H) ppm.

20

Part B. To the product from part A (1 g, 3.6 mmol) in CH₂Cl₂ (75 mL) in a separatory funnel were added cold 1N NaOH (25 mL) and 4-chlorobutylchloride (0.53 mL, 4.7 mmol). The reaction was shaken for 15 min, then separated and the

25

organic layer dried. To the crude amide in THF (30 mL) was added KOtBu (1.33 g, 10.9 mmol) at 0°C and the reaction was stirred 24h. The reaction was quenched with water, extracted with ethyl acetate, and dried to afford 1.1 g of
5 crude lactam that was carried onto the next step.

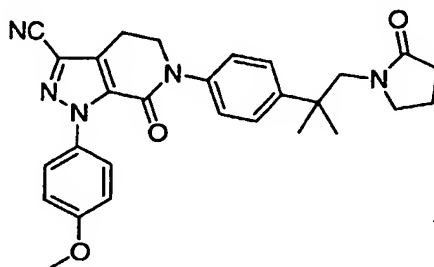
Part C. The product from part B (0.26 g, 0.76 mmol), 1-(3-chloro-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.25 g, 0.76
10 mmol), and K₂CO₃ (0.32 g, 2.3 mmol) were combined and degassed DMSO (4 mL) followed by CuI (29 mg, 0.15 mmol) were then added. The reaction was heated to 130°C for 5h. The reaction was cooled and partitioned between ethyl acetate and water and extracted with ethyl acetate and dried
15 (MgSO₄). Chromatography on silica gel using 0-5% MeOH in CH₂Cl₂ afforded ester that was carried onto the next step.

Part D. The ester from part C was placed in 5% NH₃ in ethylene glycol (1.5 mL) and heated in a sealed tube at
20 80°C for 2h. The reaction was cooled, poured into water, and filtered. Crystallization from CH₃CN/H₂O afforded 40 mg (10% for 2 steps) of the title compound. High Resolution Mass Spec for C₂₇H₂₉ClN₅O₃ (M+H)⁺ 506.1955.

25

Example 190

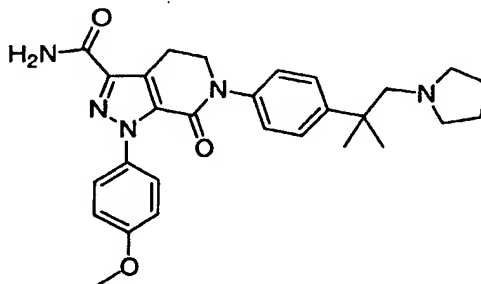
6-{4-[1,1-Dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile



To 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile (0.149 g, 0.55 mmol), 1-[2-(4-iodo-phenyl)-2-methyl-propyl]-pyrrolidin-2-one (0.19 g, 0.55 mmol), and K_2CO_3 (0.23 g, 1.7 mmol) was
5 added degassed DMSO (4 mL) followed by CuI (21 mg, 0.11 mmol). The mixture was heated to 130°C for 5h. The reaction was cooled, partitioned between ethyl acetate and water, extracted with ethyl acetate, and dried ($MgSO_4$). Chromatography on silica gel using 0-5%MeOH in CH_2Cl_2
10 followed by further purification by HPLC afforded the title compound (65 mg, 24%); HRMS for $C_{28}H_{30}N_5O_3$ ($M+H$)⁺ 484.2363.

Example 191

1-(4-Methoxy-phenyl)-6-[4-(1-methyl-1-pyrrolidin-1-ylethyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide
15



Part A. To pyrrolidine (1.2 g, 0.018 mol) in CH_2Cl_2 (50 mL) at 0°C was added 2M trimethylaluminum in heptane (9 mL, 0.018 mol) and the mixture was stirred 20 min. Ethyl-4-iodobenzoate (1 g, 3.6 mmol) was then added and the
20 reaction was stirred 72h. The reaction was quenched with ice and 1N HCl, extracted with CH_2Cl_2 , and dried ($MgSO_4$). To the crude amide was added THF (30 mL) and this solution
25 was cooled to -20°C. To this solution $TiCl_4 \cdot 2THF$ (1.2 g, 3.6 mmol) was added and stirred cold for 0.5h. A 3M diethyl ether solution of methylmagnesium bromide (7.2 mL, 21.7 mmol) was added and the reaction was stirred 24h at room temperature. Quenching with 30%NaOH, extracting with

ethyl acetate, and drying (Na_2SO_4) followed by chromatography on silica gel using 0-5%MeOH in CH_2Cl_2 afforded 1-[1-(4-iodo-phenyl)-1-methyl-ethyl]-pyrrolidine (0.1 g, 8.8%); Mass spec $(\text{M}+\text{H})^+$ 316.1.

5

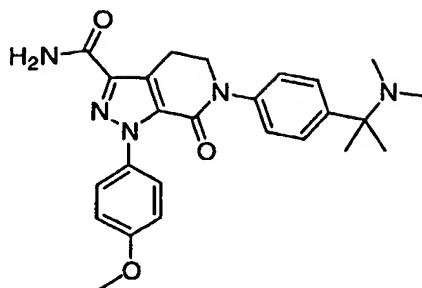
Part B. To the product from part A (100 mg, 0.32 mmol), 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (119 mg, 0.38 mmol), and K_2CO_3 (0.11 g, 0.79 mmol) was added DMSO (4 mL) and the mixture was then degassed with N_2 . CuI (12 mg, 0.063 mmol) was added. The reaction was heated to 130°C for 6h. The reaction was quenched with sat'd NaHCO_3 , extracted with CH_2Cl_2 , and dried (MgSO_4). Chromatography on silica gel using 0-5%MeOH (1% NH_3) in CH_2Cl_2 afforded 60 mg (37.7%) of ester; Mass Spec $(\text{M}+\text{H})^+$ 503.5.

Part C. To the ester (60 mg, 0.12 mmol) was added 5% NH_3 in ethylene glycol (1 mL) and the reaction was heated 80°C in a sealed tube for 2h. A solid precipitate was collected after dilution with water and the filtrate was extracted with CH_2Cl_2 . The product was purified by HPLC and freeze-dried to afford the title compound (45 mg, 64%); High Resolution Mass Spectrum for $\text{C}_{27}\text{H}_{32}\text{N}_5\text{O}_3$ $(\text{M}+\text{H})^+$ 474.2516.

25

Example 192

6-[4-(1-Dimethylamino-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide

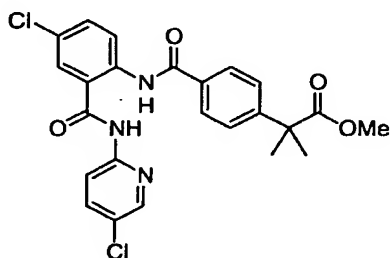


Following a procedure analogous to that used in Example 191, the title compound was prepared. High Resolution Mass Spec (M+H)⁺ for C₂₅H₃₀N₅O₃ 448.2327.

5

Example 193

2{4-[4-Chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester



10

Step A. To a solution of 2-amino-4-chloropyridine (129.0 mg, 1.0 mmol) in anhydrous THF at -78°C was added KHMDS (4.0 ml, 0.5 M solution in toluene). The mixture was stirred at this temperature under N₂ for 30 min. and a solution of 5-chloro-isatoic anhydride (198.0 mg, 1.0 mmol) in THF was added to the above mixture. The resulted mixture was warmed to rt gradually and stirred for 10 hr. The reaction mixture was quenched with sat'd NH₄Cl solution, most of the solvent was evaporated and the residue was diluted with ethyl acetate, washed with brine, and dried over MgSO₄. Removal of solvent and chromatography on silica gel (20% ethyl acetate in hexane) yielded the desired product 2-amino-5-chloro-N-(5-chloro-pyridin-2-yl)-benzamide as light brown solid. MS found: (M+1)⁺=282.2.

25

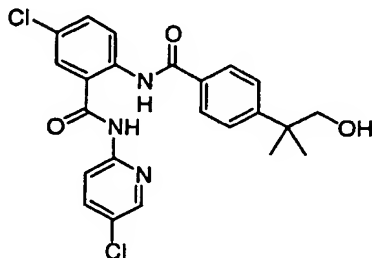
Step B. To a mixture of methyl phenylacetate (150.0 mg, 1.0 mmol) in THF at -78°C was added NaHMDS (2.2 ml, 2.2 mmol). After stirring at this temperature for 15 min, MeI (312.0 mg, 2.2 mmol) was added to the above mixture. The resulted
5 mixture was stirred at -78°C for 3 hr and rt for 1 hr. The mixture was cooled to -78°C, quenched with sat'd NH₄Cl, diluted with EtOAc, washed with aq. NaHCO₃ and brine, and dried. Flash chromatography purification (10% EtOAc in hexane) gave 2-methyl-2-phenyl-propionic acid methyl ester
10 as clear oil. MS found: (M+1)⁺=179.1.

Step C. To a suspension of AlCl₃ (500.0 mg, 3.75 mmol) in CH₂Cl₂ at -10 °C was added dropwise oxalyl chloride (476.0 mg, 3.75 mmol) in CH₂Cl₂. The mixture was stirred at this
15 temperature for 30 min. Then a solution of 2-methyl-2-phenyl-propionic acid methyl ester (178.0 mg, 1.0 mmol) in CH₂Cl₂ was added. The resulted mixture was stirred at -10 °C for 1 hr and rt overnight. The mixture was filtered through a pad of celite, the solvent and excess oxalyl
20 chloride was removed under reduced pressure. The residue was dissolved in chlorobenzene and refluxed for 4 hr. Solvent was removed and the residue was dried to give 2-(4-chlorocarbonyl-phenyl)-2-methyl-propionic acid methyl
25 ester.

Step D. To a solution of 2-(4-chlorocarbonyl-phenyl)-2-methyl-propionic acid methyl ester (240.0 mg, 1.0 mmol) in CH₂Cl₂ at 0 °C was added TEA (3.0 mmol) followed by addition of 2-amino-5-chloro-N-(5-chloro-pyridin-2-yl)-benzamide
30 (281 mg, 1.0 mmol) and DMAP (cat. 10 mg). The resulted mixture was stirred at 0 °C for 1 hr and rt over night. Solvent was evaporated and HPLC purification gave 2{4-[4-Chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester as white
35 solid. MS found: (M+1)⁺=486.2.

Example 194

2{4-[4-Chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propyl alcohol



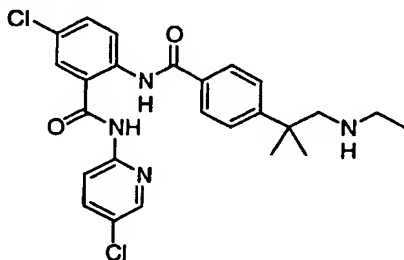
5

To a solution of 2{4-[4-Chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester (485.0 mg, 1mmol) in THF was added LiBH₄ (2.0-ml, 2.0 M solution in THF). The mixture was stirred at 60 °C over night. The reaction mixture was cooled, and quenched with sat'd NH₄Cl. HPLC purification gave 2{4-[4-Chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propyl alcohol as white solid. MS found: (M+1)⁺=457.9.

15

Example 195

5-chloro-N-(5-chloropyridin-2-yl)-2-({4-[2-(ethylamino)-1,1-dimethylethyl]benzoyl}amino)benzamide

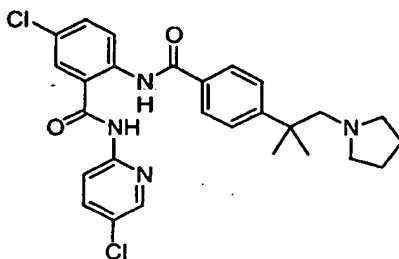


Step A. To a solution of the product obtained from Example 194 (457.0 mg, 1.0 mmol) in CH₂Cl₂ was added Dess-Martin reagent (636.0 mg, 1.5 mmol). The mixture was stirred at rt for 2.5 hr. The mixture was filtered and solvent was removed to give the desired aldehyde that was used for next step.

Step B. To a solution of the above aldehyde (46.0 mg, 0.1 mmol) in CH₂Cl₂ was added diethylamine (0.2 mmol) and NaBH₃CN (10.0 mg). The mixture was stirred at rt over
5 night. The reaction mixture was filtered and HPLC purification gave the desired product as white solid. MS found: (M+1)⁺=485.0.

Example 196

10 **5-chloro-N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)benzamide**

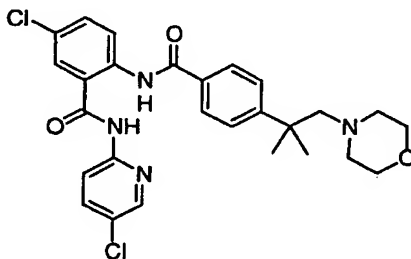


Following a procedure analogous to Example 195, the desired compound was obtained as white solid. MS found:

15 (M+1)⁺=511.3.

Example 197

5-chloro-N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl]amino)benzamide



20

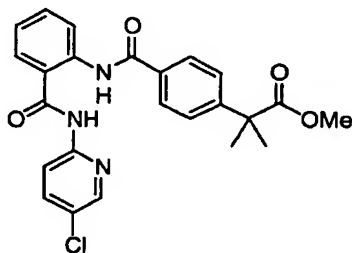
Following a procedure analogous to Example 195, the desired compound was obtained as white solid. MS found:

(M+1)⁺=527.3.

25

Example 198

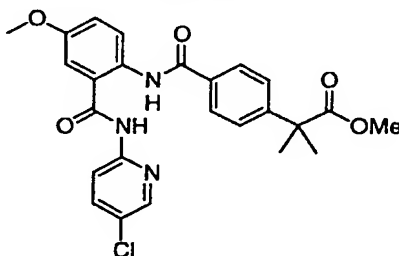
2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester



Following a procedure analogous to Example 193, the desired
5 compound was obtained as white solid. MS found:
(M+1)⁺=452.1.

Example 199

2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-4-methoxy-
10 phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl
ester

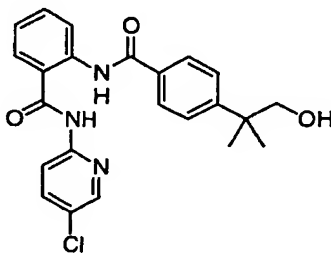


Following a procedure analogous to Example 193, the desired compound was obtained as white solid. MS found:

15 (M+1)⁺=482.1.

Example 200

***N*-(5-chloropyridin-2-yl)-2-{[4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino}benzamide**



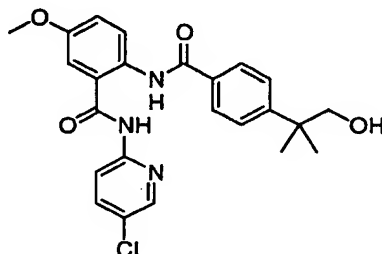
Following a procedure analogous to Example 2, the desired compound was obtained as white solid. MS found:

(M+1)⁺=424.1.

5

Example 201

N-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)-5-methoxybenzamide



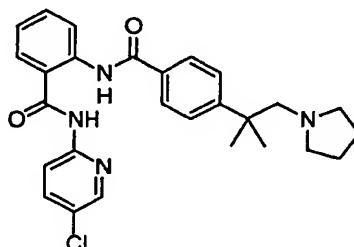
Following a procedure analogous to Example 194, the desired compound was obtained as white solid. MS found:

(M+1)⁺=454.1.

15

Example 202

N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)benzamide



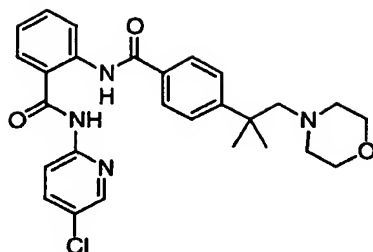
Following a procedure analogous to Example 195, the desired compound was obtained as white solid. MS found:

(M+1)⁺=577.1.

20

Example 203

N-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl]amino)benzamide



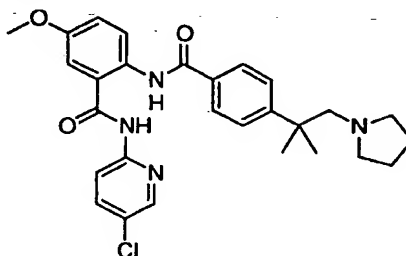
Following a procedure analogous to Example 195, the desired compound was obtained as white solid. MS found:

(M+1)⁺=493.1.

5

Example 204

N-(5-chloropyridin-2-yl)-2-((4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl)amino)-5-methoxybenzamide



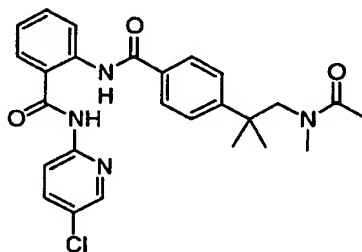
10 Following a procedure analogous to Example 195, the desired compound was obtained as white solid. MS found:

(M+1)⁺=507.1.

Example 205

15

2-((4-(2-(acetyl(methyl)amino)-1,1-dimethylethyl)benzoyl)amino)-N-(5-chloropyridin-2-yl)benzamide



Step A. Following a procedure analogous to Example 195, the
20 desired amine N-(5-chloropyridin-2-yl)-2-((4-[1,1-dimethyl-

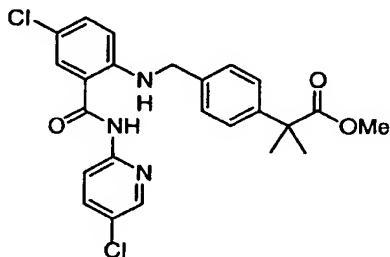
2-(methylamino)ethyl]benzoyl)-amino)benzamide was obtained as white solid.

Step B. To a solution of the above amine (10.0 mg, 0.018 mmol) in CH_2Cl_2 at 0°C was added Ac_2O (10 μl) and TEA (50 μl). The mixture was stirred at 0°C for 4 hr. Solvent was removed and the residue was purified with reverse phase HPLC. MS found: $(\text{M}+1)^+=479.2$.

10

Example 206

2-(4-([2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylamino]methyl)-phenyl)-2-methyl-propionic acid methyl ester



Step A. The product obtained from Example 193, Step C (240.0 mg, 1.0 mmol) was treated with THF/ H_2O (1:1, 10 ml). The mixture was stirred at 60°C for 2 hr. The reaction mixture was extracted with EtOAc, washed with 1N HCl, H_2O and brine. Reverse phase HPLC purification provided 4-(1-methoxycarbonyl-1-methyl-ethyl)-benzoic acid as white solid. MS found: $(\text{M}+1)^+=223.1$.

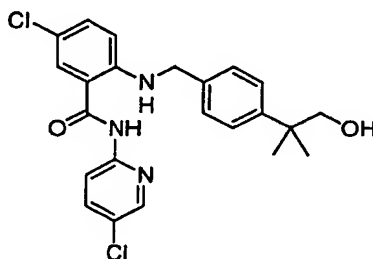
Step B. To a solution of the product obtained above (222.0 mg, 1.0 mmol) in THF was added BH_3 -THF (0.75 ml, 1.0 M solution in THF). The mixture was stirred at 65°C for 7 hr. Then the reaction mixture was cooled to rt and quenched with H_2O . After removal of solvent, the residue was purified with reverse phase to give 2-(4-hydroxymethyl-phenyl)-2-methyl-propionic acid methyl ester as clear oil. MS found: $(\text{M}+1)^+=209.2$.

Step C. To a solution of the product obtained above (83.0 mg, 0.399 mmol) in CH_2Cl_2 was added Dess-Martin reagent (203.0 mg, 0.48 mmol). The mixture was stirred at rt for 4
5 hr. The mixture was filtered, solvent was evaporated and the residue was dried to give the corresponding aldehyde, which was directly used in the next step.

Step D. A mixture of the product from above (66.0 mg, 0.32 mmol) and 2-amino-5-chloro-N-(5-chloro-pyridin-2-yl)-
10 benzamide (90.0 mg, 0.32 mmol) in ethanol was refluxed under N_2 for 2 hr. After the mixture was cooled to room temperature, NaBH_4 (100.0 mg) was added, and the resulted mixture was stirred at rt over night. The reaction mixture
15 was quenched with H_2O , and solvent was evaporated. The residue was purified with reverse phase HPLC to give desired product as white solid. MS found: $(\text{M}+1)^+=472.0$.

Example 207

20 **5-chloro-N-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzyl]amino)benzamide**

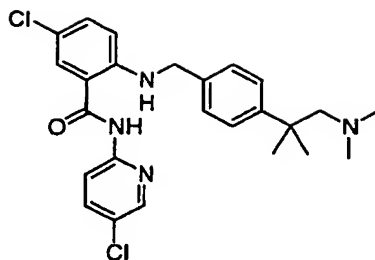


Following a procedure analogous to Example 194, the desired compound was obtained as white solid. MS found:

25 $(\text{M}+1)^+=444.1$.

Example 208

5-Chloro-N-(5-chloro-pyridin-2-yl)-2-[4-(2-dimethylamino-1,1-dimethyl-ethyl)-benzylamino]-benzamide



Step A. To a solution of 2-methyl-2-phenylpropionic acid (5.0 g, 30.49 mmol) in CH_2Cl_2 at 0°C was added oxalyl chloride (4.0 ml, 45.7 mmol). The mixture was stirred at 0°C for 3 hr. Solvent was evaporated and the residue was dried.

The above residue was dissolved in CH_2Cl_2 , and dimethylamine was purged for 20 min. or until saturated. The mixture was stirred rt for 1hr. The reaction mixture was washed with water, 1N HCl, sat'd NaHCO_3 , and brine. Chromatography purification gave N,N-dimethyl-2-phenyl-isobutyramide as white solid. MS found: $(\text{M}+1)^+=192.2$.

Step B. Following a procedure analogous to Example 193, Step C, 4-(1-dimethylcarbamoyl-1-methyl-ethyl)-benzoyl chloride was obtained as colorless oil.

A solution of the product obtained above (2.45 g, 9.7 mmol) in MeOH at 0°C was added Et_3N (40 ml) and DMAP (20 mg). The resulted mixture was stirred at 0°C for 1 hr and rt over night. Then most of the solvent was removed, the residue was diluted with EtOAc. The resulted mixture was washed with 1N HCl, water and brine. Chromatography purification (30% EtOAc in hexane) provided 4-(1-dimethylcarbamoyl-1-methyl-ethyl)-benzoic acid methyl ester as white solid. MS found: $(\text{M}+1)^+=250.1$.

Step C. To a solution of the product obtained above (35.0 mg, 0.14 mmol) in THF at 0°C was added LAH (0.7 ml, 1.0 M

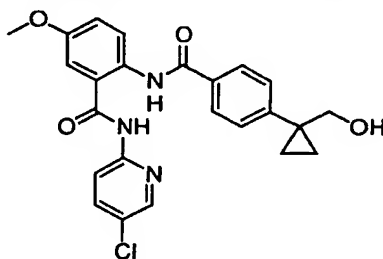
solution in THF). The mixture was stirred at 0°C for 1 hr and rt over night. Then the reaction mixture was quenched with sat'd potassium sodium tartrate solution. Solvent was evaporated and the residue was purified with reverse phase HPLC. The desired [4-(2-dimethylamino-1,1-dimethyl-ethyl)-phenyl]-methanol was obtained as clear oil. MS found: (M+1)⁺=208.2.

Step D. Following a procedure analogous to Example 195, Step A, the above alcohol was oxidized to 4-(2-dimethylamino-1,1-dimethyl-ethyl)-benzaldehyde. MS found: (M+1)⁺=206.2.

Step E. Following a procedure analogous to Example 206, Step D, the desired compound was obtained as light yellow solid. MS found: (M+1)⁺=471.1.

Example 209

N-(5-chloropyridin-2-yl)-2-((4-[1-(hydroxymethyl)cyclopropyl]benzoyl)amino)-5-methoxybenzamide



Step A. Following a procedure analogous to Example 193, Step C, 1-phenyl-cyclopropanecarboxylic acid methyl ester was converted to the desired 1-(4-chlorocarbonyl)-cyclopropanecarboxylic acid methyl ester.

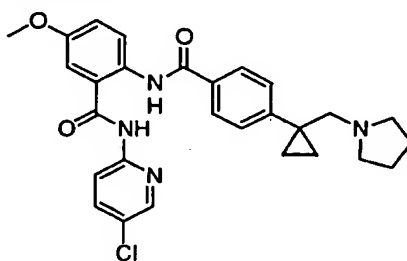
Step B. Following a procedure analogous to Example 193, Step D, the desired 1-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-4-methoxy-phenylcarbamoyl]-phenyl}-

cyclopropanecarboxylic acid methyl ester was obtained as white solid. MS found: $(M+1)^+=480.1$.

Step C. Following a procedure analogous to Example 194, the
5 desired product was obtained as yellow solid. MS found:
 $(M+1)^+=452.1$.

Example 210

N-(5-chloropyridin-2-yl)-5-methoxy-2-((4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl)amino)benzamide
10

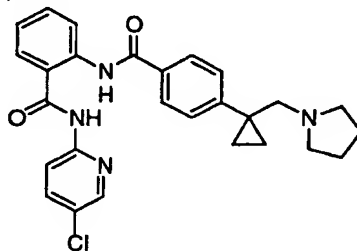


Step A. To a solution of the product obtained from Example 209 (20 mg, 0.044 mmol) in CH_2Cl_2 was added Dess-Martin reagent (28.0 mg, 0.066 mmol). The mixture was stirred at
15 rt for 2.5 hr. Then the reaction mixture was filtered, the solvent was removed and the residue was dried to give the corresponding aldehyde.

Step B. To a solution of the aldehyde from above (15.0 mg, 0.033 mmol) in 1,2-dichloroethane at 0°C was added
20 pyrrolidine (1.0 mL) and 2 drops of AcOH. The mixture was stirred at 0°C for 10 min, and $\text{NaBH}(\text{OAc})_3$ (35 mg, 0.16 mmol) was added. The resulted mixture was warmed to rt slowly and stirred for 3 hr. After quenching with H_2O , the mixture
25 was concentrated and the residue was purified with reverse phase HPLC to give the desired product as white solid. MS found: $(M+1)^+=505.2$.

Example 211

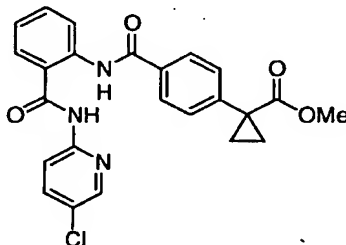
N-(5-chloropyridin-2-yl)-2-((4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl)amino)benzamide



Following a procedure analogous to Example 210, the desired
5 product was obtained as white solid. MS found:
(M+1)⁺=475.2.

Example 212

10 1-(4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-
phenyl)-cyclopropanecarboxylic acid methyl ester

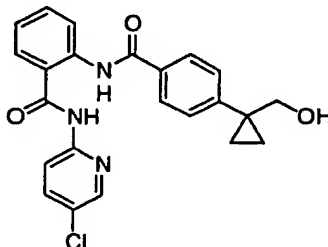


Following a procedure analogous to Example 209, Step A, the
desired product was obtained as white solid. MS found:
(M+1)⁺=450.1.

15

Example 213

N-(5-chloropyridin-2-yl)-2-((4-[1-(hydroxymethyl)cyclopropyl]benzoyl)amino)benzamide



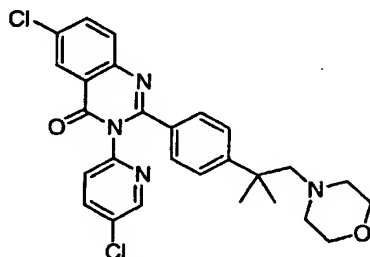
Following a procedure analogous to Example 209, the desired product was obtained as white solid. MS found:

(M+1)⁺=422.1.

5

Example 214

6-chloro-3-(5-chloropyridin-2-yl)-2-[4-(1,1-dimethyl-2-morpholin-4-ylethyl)phenyl]quinazolin-4(3H)-one

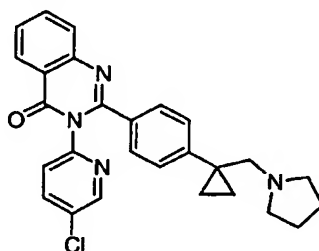


A solution of the product from Example 197 (15.0 mg, 0.028 mmol) in 5 ml of 4N HCl in dioxane and 0.5 mL of THF was refluxed for 6 hr. The mixture was cooled to rt and purified with reverse phase HPLC to give the desired product as white solid. MS found: (M+1)⁺=509.1.

15

Example 215

3-(5-chloropyridin-2-yl)-2-(4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]phenyl)quinazolin-4(3H)-one



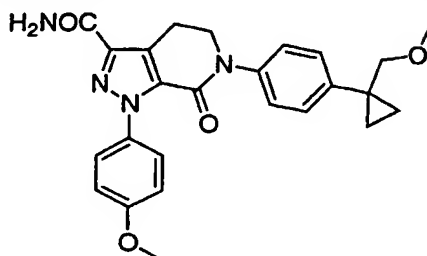
Following a procedure analogous to Example 214, the desired product was obtained as a white solid. MS found: (M+1)⁺=457.1.

25

Example 216

6-[4-(1-Methoxymethyl-cyclopropyl)-phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazo

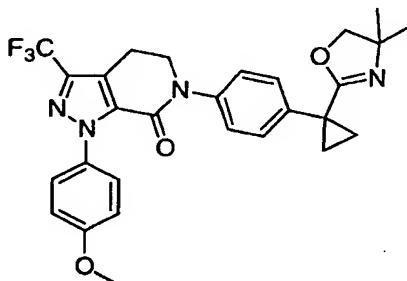
1o[3,4-c]pyridine-3-carboxylic acid amide



Following a procedure analogous to that used in Example 140, the title compound was prepared. LC/MS (ESI⁺) 447.4 (M+H).

Example 217

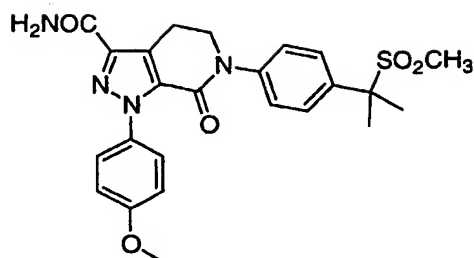
6-{4-[1-(4,4-Dimethyl-4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



Following a procedure analogous to that used in Example 140, the title compound was prepared. LC/MS (ESI⁺) 525.6 (M+H), t_R =2.05 min (10%-90% AcCN/H₂O in a 4-min run).

Example 218

6-[4-(1-Methanesulfonyl-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



- Part A. To 4-iodobenzyl bromide (5 g, 0.018 mol) in DMF (15 mL) cooled to 0°C was added sodium thiomethoxide (1.2 g, 0.017 mol). The reaction was stirred 18 h at room temperature. The reaction was partitioned between ethyl acetate and water. The aqueous layer was extracted, washed with water and brine, and dried (Na₂SO₄). The crude oil obtained was carried onto the next step.
- 10 Part B. The product of Part A (0.6 g, 2.3 mmol), 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (0.6 g, 1.9 mmol), K₂CO₃ (0.66 g, 4.7 mmol), and dimethylsulfoxide (5 mL) were combined and degassed with N₂. Copper(I) iodide (72 mg, 0.38 mmol) was added and the reaction was heated to 130°C for 5 h. The reaction was quenched with sat'd NaHCO₃, extracted with CH₂Cl₂, and dried (MgSO₄). Purification by chromatography using 1:1 hexanes/ethyl acetate afforded 0.55 g (64%) of product; High Resolution Mass Spec for
- 15 C₂₄H₂₆N₃O₄S (M+H)⁺ 452.1652.
- 20

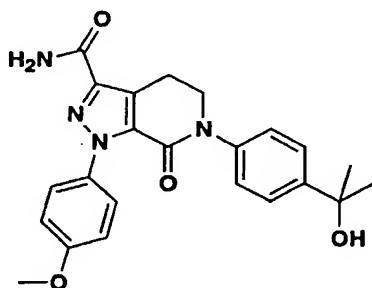
Part C. To the product of Part B (0.27 g, 0.59 mmol) in CH₂Cl₂ (15 mL) at 0°C was added 3-chloroperbenzoic acid (0.4 g) and the reaction was stirred for 72 h. The reaction was washed with sat'd NaHCO₃, and dried (MgSO₄) to afford impure product. The product was dissolved in ethyl acetate washed twice with sat'd NaHCO₃, dried (MgSO₄), filtered, and concentrated to afford 0.3 g of a yellow foam; High Resolution Mass Spec for C₂₄H₂₆N₃O₆S (M+H)⁺ 484.1541.

25

Part D. To the product of Part C (0.24 g, 4.9 mmol) in DMF (5 mL) at 0°C was added NaH (60 mg, 14.7 mmol) and iodomethane (0.09 mL, 14.7 mmol). The reaction was stirred 5 24 h, then quenched with water, extracted with ethyl acetate, and dried (MgSO₄). To the crude ester 5% NH₃ in ethylene glycol (2 mL) was added and the reaction was heated in a sealed tube at 80°C for 2h. The reaction was quenched with water and the resulting precipitate 10 collected. Purification of the solid by HPLC and freeze-drying afforded 15 mg (6%) of the title compound; High Resolution Mass Spec for C₂₄H₂₇N₄O₅S (M+H)⁺ 483.1694.

Example 219

15 **6-[4-(1-Hydroxy-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide**



Part A. To ethyl 4-iodobenzoate (1 g, 3.6 mmol) in THF (20 20 mL) at 0°C was added 3M methyl magnesium bromide (3 mL, 9 mmol). The reaction was stirred for 72h, quenched with 1N HCl, extracted with ethyl acetate, and dried (Na₂SO₄) to afford 0.94 g (100%) of the alcohol; ¹H NMR (CDCl₃) δ 7.67 (d, J=8.8Hz, 2H), 7.25 (d, J=8.8Hz, 2H), 1.56 (s, 6H) ppm.

25

Part B. The product of Part A (0.9 g, 3.4 mmol), 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (1 g, 3.4 mmol), K₂CO₃ (1.2 g, 8.5 mmol), and DMSO (10 mL) were combined and

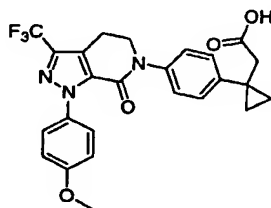
degassed with N₂. Copper(I) iodide (130 mg, 0.68 mmol) was added, and the reaction was heated to 130°C for 18h. The reaction was quenched with sat'd NaHCO₃, extracted with CH₂Cl₂, and dried (MgSO₄). Purification by chromatography
5 using 1:1 hexanes/ethyl acetate afforded an impure product; Mass Spec (M+H)⁺ 450.6.

Part C. To the impure product of Part B (0.8 g) was added 5% NH₃ in ethylene glycol (8 mL), and the reaction was
10 heated in a sealed tube at 80°C for 2h. The reaction was quenched with water and extracted with ethyl acetate. Purification of the solid by HPLC and freeze-drying afforded 120 mg of the title compound; High Resolution Mass Spec for C₂₃H₂₅N₄O₄ (M+H)⁺ 421.1862.

15

Example 220

(1-(4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-
cyclopropyl)-acetic acid



20

Part A. 1-(4-Iodophenyl)-cyclopropanecarbonyl chloride (1.74 g, 5.69 mmol) was stirred in CH₃CN and THF (1:1 v/v, 20 mL total) at 0°C under N₂. TMSCHN₂ (2M in hexanes, 4.3 mL, 1.5 eq) was added dropwise. The mixture was stirred at
25 room temperature for 4 h. It was evaporated; sat'd NaHCO₃ was added. It was extracted with EtOAc. The organic layer was washed with brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was stirred in t-BuOH (20 mL) at gentle reflux. A mixture of silver benzoate
30 (0.7 g, 3.07 mmol) and Et₃N (5 mL) was added over 1 min.

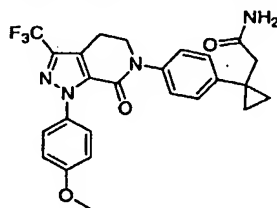
The reaction was stirred at reflux for 1 h, and the hot mixture was filtered through Celite®. H₂O was added to the filtrate; the mixture was extracted with EtOAc (3x). The organics were washed with sat'd NaHCO₃, H₂O, 1M HCl, sat'd NaHCO₃, H₂O, brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, hexanes:CH₂Cl₂ = 1:0 to 1:1 to 0:1 then 10% EtOAc in CH₂Cl₂) to give [1-(4-iodo-phenyl)-cyclopropyl]-acetic acid tert-butyl ester (0.69 g, yield: 35%). ¹H NMR (CDCl₃): δ 7.49 (d, J=8.8 Hz, 2H), 6.98 (d, J=8.8 Hz, 2H), 2.39 (s, 2H), 1.27 (s, 9H), 0.81 (s, 4H) ppm. LC/MS(ESI⁺) 359.4 (M+H).

Part B. The product from Part A (0.34 g, 0.95 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (0.31 g, 0.99 mmol) were stirred in DMSO (1 mL) in a Pyrex® tube under N₂. K₂CO₃ (0.33 g, 2.39 mmol) was added, followed by the addition of CuI (95 mg, 0.50 mmol) and 1,10-phenanthroline (90 mg, 0.50 mmol). The mixture was stirred at 120°C for 3 h. LC/MS showed 70% conversion. The cooled mixture was extracted with EtOAc (3x), washed with H₂O, brine, dried over MgSO₄, filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, hexanes:CH₂Cl₂=1:0 to 1:1 to 0:1, then EtOAc:CH₂Cl₂=1:10 to 1:2) to give (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetic acid tert-butyl ester (0.35 g, yield: 67%). ¹H NMR (CDCl₃) δ 7.48 (d, J=8.8 Hz, 2H), 7.35 (d, J=8.8 Hz, 2H), 7.21 (d, J=8.5 Hz, 2H), 6.93 (d, J=8.8 Hz, 2H), 4.12 (t, J=6.6 Hz, 2H), 3.82 (s, 3H), 3.16 (t, J=6.6 Hz, 2H), 2.51 (s, 2H), 1.39 (s, 9H), 0.91 (s, 4H) ppm. LC/MS(ESI) 542.4 (M+H).

Part C. The product from Part B (300 mg, 0.55 mmol) was stirred in CH₂CH₂ (10 mL) and TFA (5 mL) at rt for 4 h. It was purified by FCC (silica gel, EtOAc, then 10% MeOH in CH₂Cl₂) to give the desired title compound (235 mg, yield: 87.4%). ¹H NMR (CDCl₃) δ 7.43 (d, J=8.8 Hz, 2H), 7.34 (d, J=8.4 Hz, 2H), 7.19 (d, J=8.4 Hz, 2H), 6.91 (d, J=8.7 Hz, 2H), 5.23 (s, 2H), 4.11 (t, J=6.8 Hz, 2H), 3.81 (s, 3H), 3.16 (t, J=6.6 Hz, 2H), 2.63 (s, 2H), 0.95 (m, 2H), 0.91 (m, 2H) ppm. LC/MS(ESI) 486.6 (M+H). HRMS (ESI), C₂₅H₂₃N₃O₄F₃, calcd for 486.1641, found 486.1649.

Example 221

2-(1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide

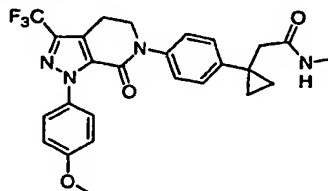


The acid chloride product from Example 210 (12.5 mg, 0.03 mmol) was stirred in THF (0.5 mL) at rt. Concentrated aqueous NH₃ (0.5 mL) was added. The mixture was stirred at rt for 4 h. LC/MS showed completion of the reaction. The mixture was purified by RP HPLC to give the title compound (9.0 mg, yield: 71.5%). ¹H NMR (CDCl₃) δ 7.38 (d, J=8.8 Hz, 2H), 7.26 (m, 4H), 6.84 (d, J=8.8 Hz, 2H), 5.23 (s, 2H), 4.04 (m, 2H), 3.74 (s, 3H), 3.08 (m, 2H), 2.48 (s, 2H), 0.91 (m, 4H) ppm. LC/MS(ESI) 485.6 (M+H).

Using the same procedure as that described for Example 221, Examples 222-224 were prepared.

Example 222

2-(1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopropyl)-N-methyl-acetamide



5

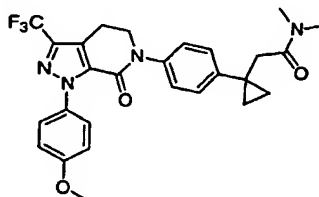
^1H NMR (CDCl_3) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.26 (m, 4H), 6.92 (d, $J=8.8$ Hz, 2H), 4.11 (t, $J=6.3$ Hz, 2H), 3.81 (s, 3H), 3.15 (t, $J=6.3$ Hz, 2H), 2.69 (m, 3H), 2.52 (s, 2H), 1.78 (m, 4H), 0.95 (m, 4H) ppm. HRMS (ESI) calcd. 499.1958; found 499.1970 for $\text{C}_{26}\text{H}_{26}\text{F}_3\text{N}_4\text{O}_3$ (M+H). LC/MS (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.30$ min), 499.6 (M+H).

10

Example 223

2-(1-{4-[1-(4-Methoxy-phenyl)-7-oxo-3-trifluoromethyl-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopropyl)-N,N-dimethyl-acetamide

15



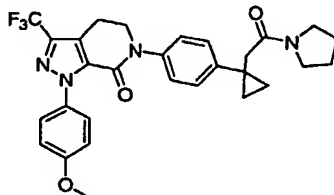
^1H NMR (CDCl_3) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.38 (d, $J=8.5$ Hz, 2H), 7.19 (d, $J=8.4$ Hz, 2H), 6.91 (d, $J=8.8$ Hz, 2H), 4.10 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.13 (t, $J=6.6$ Hz, 2H), 2.86 (s, 3H), 2.76 (s, 3H), 2.64 (s, 3H), 0.89 (m, 4H) ppm. HRMS (ESI), calcd. 513.2114; found 513.2113 for $\text{C}_{27}\text{H}_{28}\text{F}_3\text{N}_4\text{O}_3$ (M+H). LC/MS (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.46$ min), 513.6 (M+H).

20

25

Example 224

1-(4-Methoxy-phenyl)-6-{4-[1-(2-oxo-2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



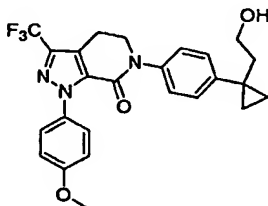
5

^1H NMR (CDCl_3) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.41 (d, $J=8.4$ Hz, 2H), 7.18 (d, $J=8.4$ Hz, 2H), 6.91 (d, $J=8.8$ Hz, 2H), 4.10 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.38 (t, $J=6.4$ Hz, 2H), 3.13 (m, 4H), 2.56 (s, 2H), 1.78 (m, 4H), 0.92 (m, 2H), 0.88 (m, 2H) ppm. HRMS(ESI) calcd. 539.2271; found 539.2214 for $\text{C}_{28}\text{H}_{30}\text{F}_3\text{N}_3\text{O}_3$ ($\text{M}+\text{H}$). LC/MS (ESI) (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.52$ min) 539.6 ($\text{M}+\text{H}$).

10

Example 225

6-{4-[1-(2-Hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



15

Using the similar sequence for the preparation of Part E in Example 1 but using the product of Example 220 as the starting material, the title compound was prepared. ^1H NMR (CDCl_3) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.31 (d, $J=8.8$ Hz, 2H), 7.21 (d, $J=8.5$ Hz, 2H), 6.92 (d, $J=9.2$ Hz, 2H), 5.23 (s, 2H), 4.11 (t, $J=6.6$ Hz, 2H), 3.80 (s, 3H), 3.59 (t, $J=7.0$ Hz, 2H), 3.14 (t, $J=6.6$ Hz, 2H), 1.82 (t, $J=7.0$ Hz, 2H), 0.78 (m, 2H), 0.74 (m, 2H) ppm. LC/MS (ESI) 472.4 ($\text{M}+\text{H}$).

25

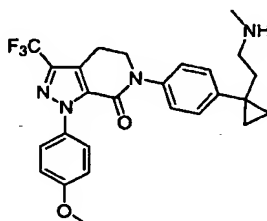
Following procedures analogous to that used for Part E and Part F of Example 1, but using the product of Example 225 as one of the starting materials, Examples 226-229 were prepared.

5

Example 226

1-(4-Methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

10



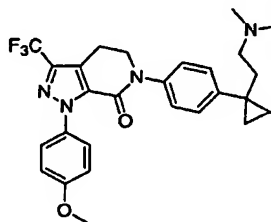
^1H NMR (CDCl_3) δ 7.45 (d, $J=8.8$ Hz, 2H), 7.26 (m, 4H), 6.91 (d, $J=8.8$ Hz, 2H), 4.12 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.16 (t, $J=6.6$ Hz, 2H), 2.98 (m, 2H), 2.75 (m, 6H), 1.97 (m, 2H), 0.86 (m, 2H), 0.79 (m, 2H) ppm. HRMS (ESI) calcd. 485.2165; found 485.2153 for $\text{C}_{26}\text{H}_{28}\text{F}_3\text{N}_4\text{O}_2$ (M+H). LC/MS (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.10$ min) 485.6 (M+H).

15

Example 227

6-{4-[1-(2-Dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt

20



^1H NMR (CDCl_3) δ 7.45 (d, $J=8.8$ Hz, 2H), 7.26 (m, 4H), 6.91 (d, $J=8.8$ Hz, 2H), 4.12 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.16 (t, $J=6.6$ Hz, 2H), 2.98 (m, 2H), 2.75 (m, 6H), 1.97

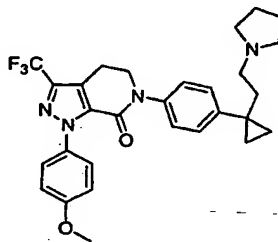
25

(m, 2H), 0.86 (m, 2H), 0.79 (m, 2H) ppm. HRMS (ESI) calcd. 499.2322; found 499.2318 for $C_{27}H_{30}F_3N_4O_2$ (M+H). LC/MS (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.10$ min) 499.6 (M+H).

5

Example 228

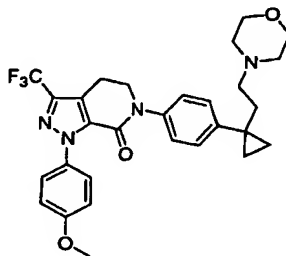
1-(4-Methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydropyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



10 1H NMR ($CDCl_3$) δ 7.46 (d, $J=8.8$ Hz, 2H), 7.26 (m, 4H), 6.92 (d, $J=8.8$ Hz, 2H), 4.12 (t, $J=6.6$ Hz, 2H), 3.81 (s, 3H), 3.77 (m, 2H), 3.16 (t, $J=6.6$ Hz, 2H), 3.02 (m, 2H), 2.65 (m, 2H), 2.02 (m, 6H), 0.84 (m, 2H), 0.77 (m, 2H).
HRMS(ESI) calcd. 525.2478, found 525.2483 for $C_{29}H_{32}F_3N_4O_2$ (M+H).
15 LC/MS (10-90% CH_3CN in H_2O in 4-min run, $t_R=2.18$ min) 525.6 (M+H).

Example 229

1-(4-Methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydropyrazolo[3,4-c]pyridin-7-one, trifluoroacetic acid salt



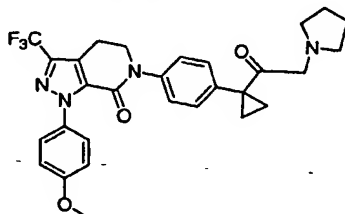
1H NMR ($CDCl_3$) δ 7.45 (d, $J=8.8$ Hz, 2H), 7.26 (m, 4H), 6.92 (d, $J=8.8$ Hz, 2H), 4.12 (t, $J=6.6$ Hz, 2H), 3.93 (m, 4H),
25 3.81 (s, 3H), 3.47 (m, 2H), 3.16 (t, $J=6.6$ Hz, 2H), 2.98

(m, 2H), 2.72 (m, 2H), 1.99 (m, 2H), 0.85 (m, 2H), 0.76 (m, 2H) ppm. HRMS(ESI) calcd. 541.2427, found 541.2413 for $C_{29}H_{32}F_3N_4O_3$ (M+H). LC/MS (ESI) (10-90% CH_3CN in H_2O in a 4-min run, $t_R=2.11$ min), 541.6 (M+H).

5

Example 230

1-(4-Methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-acetyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one



10

Part A. 1-(4-Iodophenyl)-cyclopropanecarbonyl chloride (1.74 g, 5.69 mmol) was stirred in CH_3CN and THF (1:1 v/v, 20 mL) total at 0°C under N_2 . $TMSCHN_2$ (2M in hexanes, 4.3 mL) was added dropwise. The mixture was stirred at 0°C to rt for 2 h. It was partitioned between EtOAc and sat'd $NaHCO_3$. The organics were washed with brine, dried over $MgSO_4$, filtered, and concentrated to dryness. $HBr/HOAc$ (30%, 8 mL) was added dropwise to the residue at 0°C. The mixture was stirred at 0°C for 30 min, and EtOAc was added; it was washed with 15% citric acid, sat'd $NaHCO_3$, H_2O , brine, dried over $MgSO_4$, filtered, and concentrated to dryness to give crude 2-bromo-1-[1-(4-iodo-phenyl)-cyclopropyl]-ethanone (0.5 g).

Part B. The product from Part A (0.5 g, 1.37 mmol) was dissolved in DMF (1.8 mL), a spatula tip of the K_2CO_3 and pyrrolidine (0.2 mL) were added. The mixture was heated at 80°C for 1.5 h. The cooled mixture was partitioned between EtOAc and H_2O . The organics were washed with brine, dried over $MgSO_4$, filtered, and concentrated to dryness. The

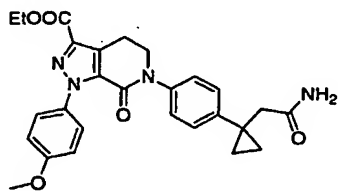
30

residue was purified by FCC (silica gel, EtOAc:CH₂Cl₂=0:1 to 1:0, then 10% MeOH in EtOAc) to give 1-[1-(4-iodo-phenyl)-cyclopropyl]-2-pyrrolidin-1-yl-ethanone (85 mg, yield: 17% for 2 steps). ¹H NMR (CDCl₃) δ 7.68 (d, J=8.4 Hz, 2H), 7.13 (d, J=8.4 Hz, 2H), 3.30 (s, 2H), 2.51 (m, 4H), 1.75 (m, 4H), 1.63 (m, 2H) 1.10 (m, 2H) ppm. LC/MS (ESI) 356.4 (M+H).

Part C. The product of part B (50 mg, 0.14 mmol) and 1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one (52.5 mg, 0.17 mmol) were stirred in DMSO (0.2 mL) in a Pyrex tube under N₂. K₂CO₃ (39 mg) was added followed by the addition of CuI (20 mg) and 9,10-phenanthroline (20 mg). The mixture was stirred at 120°C for 2 h. The cooled mixture was purified by reverse phase HPLC to give the title compound (9.6 mg, yield: 12.7 %). LC/MS (ESI) 539.6 (M+H).

Example 231

6-[4-(1-Carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester



Part A. Following procedure similar to that of Part C in Example 220 but using 1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester and [1-(4-iodo-phenyl)-cyclopropyl]-acetic acid tert-butyl ester as starting materials, 6-[4-(1-tert-butoxycarbonylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester was obtained (406.29

mg, yield: 86%). ^1H NMR (CDCl_3) δ 7.48 (d, $J=8.9$ Hz, 2H), 7.33 (d, $J=8.5$ Hz, 2H), 7.22 (d, $J=8.5$ Hz, 2H), 6.91 (d, $J=8.9$ Hz, 2H), 4.47 (q, $J=7.1$ Hz, 2H), 4.12 (m, 2H), 3.81 (s, 3H), 3.32 (t, $J=6.6$ Hz, 2H), 2.50 (s, 2H), 1.44 (t, $J=7.1$ Hz, 3H), 1.38 (s, 9H), 0.90 (s, 4H) ppm. LRMS (ESI) 546.2 (M+H).

Part B. The product from Part A (450 mg) was stirred in a mixture of CH_2Cl_2 (10 mL) and TFA (15 mL) at rt for 4h. The solvents were evaporated. The residue was dried in vacuo to give 6-[4-(1-carboxymethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester (386 mg, yield: 95.6%). ^1H NMR (CDCl_3) δ 7.45 (d, $J=8.7$ Hz, 2H), 7.35 (d, $J=8.3$ Hz, 2H), 7.21 (d, $J=8.3$ Hz, 2H), 6.91 (d, $J=8.7$ Hz, 2H), 4.47 (q, $J=7.1$ Hz, 2H), 4.14 (t, $J=6.6$ Hz, 2H), 3.82 (s, 3H), 3.34 (t, $J=6.5$ Hz, 2H), 2.65 (s, 2H), 1.43 (t, $J=7.1$ Hz, 3H), 0.97 (m, 2H), 0.91 (m, 2H) ppm. LRMS (ESI) 490.1 (M+H).

Part C. The product from Part B (150 mg, 0.31 mmol) was stirred in CH_2Cl_2 (5 mL). Oxalyl chloride (2M solution in CH_2Cl_2 , 0.3 mL. ca. 2 eq) was added, followed by the addition of 1 drop of DMF. The mixture was stirred at rt for 1h. The solvents were evaporated. The residue was dried in vacuo. One third of the residue (0.1 mmol) was dissolved in THF (2.0 mL), concentrated $\text{NH}_3\cdot\text{H}_2\text{O}$ (2.0 mL) was added. The mixture was stirred at rt for 2h. EtOAc was added. It was washed with H_2O , brine, dried over MgSO_4 , filtered, and concentrated to dryness. The residue was purified by FCC (silica gel, CH_2Cl_2 , then EtOAc) to give pure title compound. ^1H NMR (CDCl_3) δ 7.39 (d, $J=8.9$ Hz, 2H), 7.22 (d, $J=8.4$ Hz, 2H), 7.15 (d, $J=8.5$ Hz, 2H), 6.82 (d, $J=8.9$ Hz, 2H), 5.46 (s, br, 1H), 5.30 (s, br, 1H), 4.38

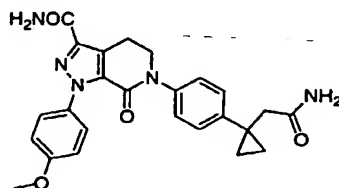
(t, $J=7.1$ Hz, 2H), 4.03 (t, $J=6.6$ Hz, 2H), 3.73 (s, 3H), 3.23 (t, $J=6.7$ Hz, 2H), 2.45 (s, 2H), 1.36 (t, $J=7.1$ Hz, 2H), 0.88 (m, 4H) ppm. HRMS (ESI) $C_{27}H_{29}N_4O_5$ calcd for 489.2138, found 489.2152.

5

Following procedures analogous to that used for Example 231, Examples 232-238 were prepared.

Example 232

10 6-[4-(1-Carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide

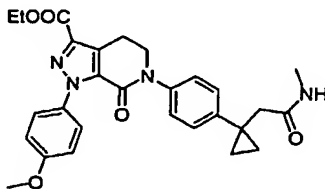


1H NMR (methanol- d_4) δ 7.50 (d, $J=8.9$ Hz, 2H), 7.40 (d, $J=8.5$ Hz, 2H), 7.27 (d, $J=8.5$ Hz, 2H), 7.00 (d, $J=8.9$ Hz, 2H), 4.10 (t, $J=6.6$ Hz, 2H), 3.86 (s, 3H), 3.33 (m, 2H), 2.54 (s, 2H), 0.99 (m, 2H), 0.94 (m, 2H) ppm. LC/MS (ESI) $t_R=2.66$ min (10-90% MeOH in H_2O with 10 mM NH_4OAc in a 4-min gradient), 460.10 (M+H, 100%), 492.11 (M+H+MeOH, 70%).

20

Example 233

1-(4-Methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester



25

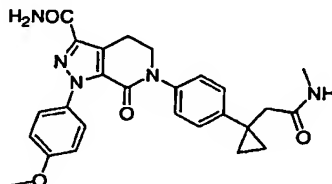
1H NMR ($CDCl_3$) δ 7.39 (d, $J=9.0$ Hz, 2H), 7.17 (m, 4H), 6.82 (d, $J=9.0$ Hz, 2H), 5.46 (m, 1H), 4.38 (q, $J=6.8$ Hz, 2H), 4.02 (t, $J=6.8$ Hz, 2H), 3.73 (s, 3H), 3.23 (t, $J=6.6$ Hz,

2H), 2.61 (d, $J=4.8$ Hz, 3H), 2.43 (s, 2H), 1.36 (t, $J=7.2$ Hz, 3H), 0.87 (s, 4H) ppm. HRMS (ESI) $C_{28}H_{31}N_4O_5$, calcd for 503.2294, found 503.2281 .

5

Example 234

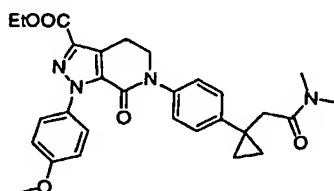
1-(4-Methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



- 10 1H NMR (methanol- d_4) δ 7.38 (d, $J=8.8$ Hz, -2H), 7.23 (d, $J=8.8$ Hz, 2H), 7.12 (d, $J=8.8$ Hz, 2H), 6.85 (d, $J=8.8$ Hz, 2H), 3.94 (t, $J=6.6$ Hz, 2H), 3.71 (s, 3H), 3.18 (t, $J=6.6$ Hz, 2H), 2.48 (s, 3H), 2.35 (s, 2H), 0.84 (m, 2H), 0.77 (m, 2H) ppm. LC/MS (ESI) $t_R=2.76$ min (10-90% MeOH in H_2O with
- 15 10 mM NH_4OAc in a 4-min gradient), 474.09 (M+H, 100%), 506.12 (M+H+MeOH, 100%).

Example 235

- 6-[4-(1-Dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester
- 20

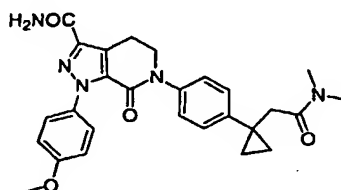


- 1H NMR ($CDCl_3$) δ 7.40 (d, $J=8.9$ Hz, 2H), 7.30 (d, $J=8.4$ Hz, 2H), 7.13 (d, $J=8.4$ Hz, 2H), 6.82 (d, $J=9.0$ Hz, 2H), 4.38 (q, $J=7.1$ Hz, 2H), 4.03 (t, $J=6.6$ Hz, 2H), 3.73 (s, 3H), 3.22 (t, $J=6.6$ Hz, 2H), 2.78 (s, 3H), 2.68 (s, 3H), 2.57 (s, 2H), 1.36 (t, $J=7.1$ Hz, 3H), 0.82 (m, 2H), 0.81 (m, 2H)
- 25

ppm. HRMS (ESI) $C_{29}H_{33}N_4O_5$, calcd for 517.2451, found 517.2439 .

Example 236

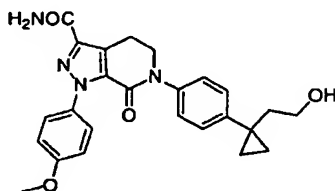
5 6-[4-(1-Dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



1H NMR (methanol- d_4) δ 7.50 (d, $J=9.0$ Hz, 2H), 7.38 (d, $J=8.5$ Hz, 2H), 7.26 (d, $J=8.5$ Hz, 2H), 6.99 (d, $J=9.0$ Hz, 2H), 4.08 (t, $J=6.6$ Hz, 2H), 3.85 (s, 3H), 3.33 (m, 2H), 2.83 (s, 3H), 2.75 (s, 3H), 2.72 (s, 2H), 0.95 (m, 2H), 0.90 (m, 2H) ppm. LC/MS (ESI) $t_R=2.90$ min (10-90% MeOH in H_2O with 10 mM NH_4OAc in a 4-min gradient), 488.10 (M+H, 100%), 520.13 (M+H+MeOH, 60%).

Example 237

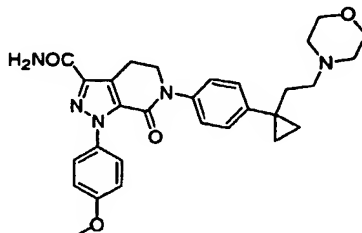
20 6-[4-[1-(2-Hydroxy-ethyl)-cyclopropyl]-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



1H NMR (methanol- d_4) δ 7.43 (d, $J=8.8$ Hz, 2H), 7.29 (d, $J=8.4$ Hz, 2H), 7.21 (d, $J=8.5$ Hz, 2H), 6.92 (d, $J=8.9$ Hz, 2H), 4.03 (m, 4H), 3.78 (s, 3H), 3.43 (t, $J=6.6$ Hz, 2H), 1.78 (t, $J=7.0$ Hz, 2H), 0.72 (m, 2H), 0.70 (m, 2H) ppm. LC/MS (ESI) $t_R=2.62$ min (10-90% MeOH in H_2O with 10 mM NH_4OAc in a 4-min gradient), 474.11 (M+H, 100%).

Example 238

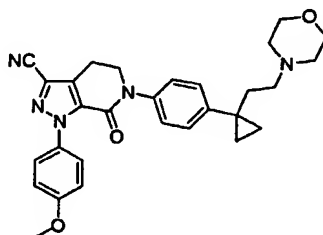
1-(4-Methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-
cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide



¹H NMR (methanol-*d*₄) δ 7.38 (d, *J*=9.0 Hz, 2H), 7.24 (d, *J*=8.5 Hz, 2H), 7.15 (d, *J*=8.5 Hz, 2H), 6.87 (d, *J*=9.0 Hz, 2H), 3.97 (t, *J*=6.6 Hz, 2H), 3.73 (s, 3H), 3.52 (m, 4H), 3.20 (m, 2H), 2.25 (m, 6H), 1.70 (m, 2H), 0.70 (m, 2H), 0.65 (m, 2H) ppm. LC/MS (ESI) *t*_R=3.18 min (10-90% MeOH in H₂O with 10 mM NH₄OAc in a 4-min gradient), 516.13 (M+H, 100%).

Example 239

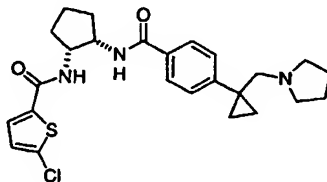
1-(4-Methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-
cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile, trifluoroacetic
acid salt



Following the procedure as Example 74 but using the product of Example 238 as the starting material, the titled compound was prepared. LRMS (ESI) 498.2 (M+H).

Example 240

(1*R*, 2*S*)-5-Chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide, trifluoroacetic acid salt



5

Part A. To a solution of (1*S*, 2*S*)-2-benzyloxy-cyclopentylamine (9.8 g, 51.2 mmol) in THF (150 mL) were sequentially added Et₃N (13.6 mL, 0.10 mol) and (Boc)₂O (12.30 g, 56.4 mmol) at 0°C. The reaction mixture was stirred overnight at room temperature, and diluted with EtOAc (200 mL). The organic phase was washed with H₂O, brine, and dried (Na₂SO₄). The solvent was evaporated to afford (1*S*, 2*S*)-(2-benzyloxy-cyclopentyl)-carbamic acid tert-butyl ester (14.90 g, 100%) as a slight yellow solid.

15 MS *m/z* 293.0 ([*M* + *H*]⁺).

Part B. The product from Part A (10.0 mg, 34.2 mmol) was dissolved in ethanol (100 mL), Pd/C (800 mg, 5%) was then added. The reaction mixture was hydrogenated at 25 psi with stirring for 4 h, and filtered through a pad of Celite®. The filtrate was evaporated to afford (1*S*, 2*S*)-(2-hydroxy-cyclopentyl)-carbamic acid tert-butyl ester (6.84 g, 99%) as a white solid. MS *m/z* 202.0 ([*M* + *H*]⁺).

20

Part C. To a solution of the product from Part B (4.95 g, 24.6 mmol) in CH₂Cl₂ (50 mL) were sequentially added Et₃N (4.11 mL, 29.51 mol) and MsCl (2.09 g, 27.05 mmol) at 0°C. The reaction mixture was stirred for 2 h at 0 °C, then quenched with H₂O and extracted with EtOAc (3 x 50 mL).

30 The organic phase was washed with H₂O, brine, and dried (Na₂SO₄). The solvent was evaporated to afford (1*S*, 2*S*)-

methanesulfonic acid 2-tert-butoxycarbonylamino-cyclopentyl ester (6.35 g, 92%) as a white solid. MS m/z 297.0 ($[M + NH_4]^+$).

5 Part D. NaN_3 (4.40 g, 67.7 mmol) was added to a solution of the product from Part C (6.30 g, 22.6 mmol) in DMF (50 mL), and the reaction mixture was heated at 80 °C for 12 h with vigorous stirring. The reaction was cooled to room temperature, poured into water, and extracted with EtOAc (4
10 x 100 mL). The extracts were combined and washed with H_2O , aqueous LiCl (10%), brine, and dried (Na_2SO_4). The solvent was evaporated, and the residue was taken to next step without purification. The residue from above reaction was then dissolved in ethanol (200 mL), and Pd/C (300 mg, 5%)
15 was added. The reaction mixture was hydrogenated at 1 atm with stirring for 24 h, and filtered through a pad of Celite®. The filtrate was evaporated to afford (1S, 2R)-(2-amino-cyclopentyl)-carbamic acid tert-butyl ester (6.84 g, 99%) as a white solid. MS m/z 201.0 ($[M + H]^+$).

20 Part E. The product from Part D (150 mg, 0.75 mmol) and 5-chloro-thiophene-2-carboxylic acid (101 mg, 0.62 mmol) were dissolved in DMF (2 mL) and cooled to 0°C. To this solution was added HATU (354 mg, 0.93 mmol), DIEA (0.22 mL, 1.24
25 mmol). The mixture was stirred overnight. It was diluted with ethyl acetate, washed with water, aqueous LiCl (10%), brine, and dried ($MgSO_4$). After evaporation of the solvent, the residue was purified on silica gel using 50% EtOAc-Hexane to afford (1S, 2R)-(2-[(5-chloro-thiophene-2-carbonyl)-amino]-cyclopentyl)-carbamic acid tert-butyl
30 ester (115 mg, 54%) as a white solid. MS m/z 367.6 ($[M+Na]^+$).

Part F. The product from Part E (115 mg, 0.33 mmol) was
35 suspended in CH_2Cl_2 (1 mL) and TFA (1 mL) was added. A

clear solution was obtained and stirred for 2 h at ambient temperature. The resulting solution was concentrated, and the residue was partitioned between EtOAc and aqueous Na_2CO_3 . The aqueous was extracted with EtOAc (3 x 10 mL).

- 5 The extracts were combined and washed with brine and dried (Na_2SO_4). Evaporation of the solvent afforded (1*R*, 2*S*)-5-chloro-thiophene-2-carboxylic acid (2-amino-cyclopentyl)-amide (80 mg, 98%) as a white solid that was taken to next step without purification. MS m/z 245.0 ($[\text{M} + \text{H}]^+$).

10

Part G. The product from Part F (40 mg, 0.16 mmol) and excess 4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl chloride (Example 1) and DIEA (0.05 mL) were stirred in CH_2Cl_2 (1 mL) at 0°C. The above solution was added Et_3N

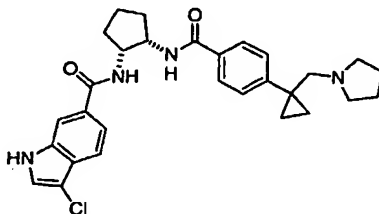
- 15 (0.05 mL) and stirred overnight. The reaction mixture was diluted with ethyl acetate, washed with water, aqueous LiCl (10%), brine, and dried (MgSO_4). After evaporation of the solvent, the residue was purified on reverse-phase HPLC to afford the title compound as a white solid. LRMS (ESI)
20 472.2 ($\text{M}+\text{H}$).

Using the same procedure as that described for Example 240, Examples 241-243 were prepared:

25

Example 241

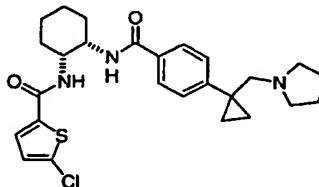
(1*R*, 2*S*)-3-Chloro-1*H*-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide, trifluoroacetic acid salt



- 30 LC/MS (ESI) 505.2 ($\text{M}+\text{H}$).

Example 242

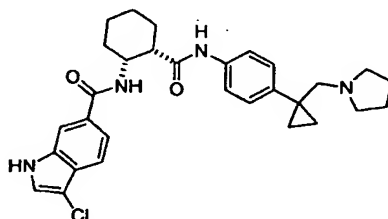
(1*R*, 2*S*)-5-Chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide, trifluoroacetic acid salt



LC/MS (ESI) 486.2 (M+H)

Example 243

Cis-3-Chloro-1*H*-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-phenylcarbamoyl]-cyclohexyl}-amide, trifluoroacetic acid salt

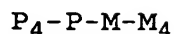


LRMS (ESI) 519.2 (M+H).

Numerous modifications and variations of the present invention are possible in light of the above teachings. It is therefore to be understood that within the scope of the appended claims, the invention may be practiced otherwise than as specifically described herein.

WHAT IS CLAIMED IS:

1. A compound of formula I:



I

5 or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

M is a 3-10 membered carbocycle or a 4-10 membered heterocycle, consisting of: carbon atoms and 1-3
10 heteroatoms selected from O, S(O)_p, N, and NZ²;

ring M is substituted with 0-3 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

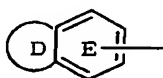
15 P is fused onto ring M and is a 5, 6, or 7 membered carbocycle or a 5, 6, or 7 membered heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

20 ring P is substituted with 0-3 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

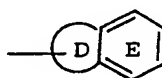
alternatively, ring P is absent and P₄ is directly attached to ring M, provided that when ring P is absent, P₄ and
25 M₄ are attached to the 1,2, 1,3, or 1,4 positions of ring M;

one of P₄ and M₄ is -Z-A-B and the other -G₁-G, provided that P₄ and M₄ are attached to different rings when
30 ring P is present;

G is a group of formula IIa or IIb:



IIa



IIb

ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

5

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;

E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

10

alternatively, ring D is absent and ring E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, pyrrolyl, pyrazolyl, imidazolyl, isoxazolyl, oxazolyl, triazolyl, thienyl, and thiazolyl, and ring E is substituted with 1-3 R;

15

alternatively, ring D is absent and ring E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, pyrrolyl, pyrazolyl, imidazolyl, isoxazolyl, oxazolyl, triazolyl, thienyl, and thiazolyl, and ring E is substituted with 1 R and with a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, wherein the 5-6 membered heterocycle is substituted with 0-2 carbonyls and 1-3 R and there are 0-3 ring double bonds;

20

25

R is selected from H, C₁₋₄ alkyl, F, Cl, Br, I, OH, OCH₃, OCH₂CH₃, OCH(CH₃)₂, OCH₂CH₂CH₃, CN, C(=NR⁸)NR⁷R⁹, NHC(=NR⁸)NR⁷R⁹, ONHC(=NR⁸)NR⁷R⁹, NR⁸CH(=NR⁷), NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂, CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, CH₂CH₂NH₂, CH₂CH₂NH(C₁₋₃ alkyl), CH₂CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tC(O)H, (CR⁸R⁹)_tC(O)R^{2c}, (CR⁸R⁹)_tNR⁷R⁸,

30

35

$(\text{CR}^8\text{R}^9)_t\text{C}(\text{O})\text{NR}^7\text{R}^8$, $(\text{CR}^8\text{R}^9)_t\text{NR}^7\text{C}(\text{O})\text{R}^7$, $(\text{CR}^8\text{R}^9)_t\text{OR}^3$,
 $(\text{CR}^8\text{R}^9)_t\text{S}(\text{O})_p\text{NR}^7\text{R}^8$, $(\text{CR}^8\text{R}^9)_t\text{NR}^7\text{S}(\text{O})_p\text{R}^7$, $(\text{CR}^8\text{R}^9)_t\text{SR}^3$,
 $(\text{CR}^8\text{R}^9)_t\text{S}(\text{O})\text{R}^3$, $(\text{CR}^8\text{R}^9)_t\text{S}(\text{O})_2\text{R}^3$, and OCF_3 , provided that
 $\text{S}(\text{O})_p\text{R}^7$ forms other than $\text{S}(\text{O})_2\text{H}$ or $\text{S}(\text{O})\text{H}$;

5

alternatively, when 2 R groups are attached to adjacent
 atoms, they combine to form methylenedioxy or
 ethylenedioxy;

10 A is selected from:

C_3 -10 carbocycle substituted with 0-2 R^4 , and

5-12 membered heterocycle substituted with 0-2 R^4 and
 consisting of: carbon atoms and 1-4 heteroatoms selected
 from the group consisting of N, O, and $\text{S}(\text{O})_p$;

15

B is Y-R^{4a} or X-Y-R^{4a} , provided that Z and B are attached to
 different atoms on A and A and R^{4a} or X and R^{4a} are
 attached to the same atom on Y;

20 X is selected from $-(\text{CR}^2\text{R}^{2a})_{1-4}-$, $-\text{CR}^2(\text{CR}^2\text{R}^{2b})(\text{CH}_2)_t-$,
 $-\text{C}(\text{O})-$, $-\text{C}(=\text{NR}^{1b})-$, $-\text{CR}^2(\text{NR}^{1b}\text{R}^2)-$, $-\text{CR}^2(\text{OR}^2)-$,
 $-\text{CR}^2(\text{SR}^2)-$, $-\text{C}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{C}(\text{O})-$, $-\text{S}(\text{O})-$, $-\text{S}(\text{O})_2-$,
 $-\text{SCR}^2\text{R}^{2a}-$, $-\text{S}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{S}(\text{O})_2\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{S}-$,
 $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})-$, $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})_2-$, $-\text{S}(\text{O})_2\text{NR}^2-$,
 25 $-\text{S}(\text{O})_2\text{NR}^2\text{CR}^2\text{R}^{2a}-$, $-\text{CR}^2\text{R}^{2a}\text{S}(\text{O})_2\text{NR}^2-$, $-\text{NR}^2\text{S}(\text{O})_2-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2\text{S}(\text{O})_2-$, $-\text{NR}^2\text{S}(\text{O})_2\text{CR}^2\text{R}^{2a}-$, $-\text{NR}^2\text{C}(\text{O})-$,
 $-\text{C}(\text{O})\text{NR}^2-$, $-\text{NR}^2\text{C}(\text{O})\text{CR}^2\text{R}^{2a}-$, $-\text{C}(\text{O})\text{NR}^2\text{CR}^2\text{R}^{2a}-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2\text{C}(\text{O})-$, $-\text{CR}^2\text{R}^{2a}\text{C}(\text{O})\text{NR}^2-$, NR^2 , $-\text{NR}^2\text{CR}^2\text{R}^{2a}-$,
 $-\text{CR}^2\text{R}^{2a}\text{NR}^2-$, O, $-\text{OCR}^2\text{R}^{2a}-$, and $-\text{CR}^2\text{R}^{2a}\text{O}-$;

30

Y is a C_3 -10 carbocycle or 3-10 membered heterocycle,
 wherein the carbocycle or heterocycle consists of
 carbon atoms and 0-4 heteroatoms selected from N, O,
 and $\text{S}(\text{O})_p$, the carbocycle or heterocycle further

comprises 0-4 double bonds and 0-2 carbonyl groups,
and the carbocycle or heterocycle is substituted with
0-2 R⁴, provided that Y is other than a 1,3-dioxolanyl
group;

5

alternatively, Y is CY¹Y², and Y¹ and Y² are independently
C₁₋₄ alkyl substituted with 0-2 R⁴;

G₁ is absent or is selected from (CR³R^{3a})₁₋₅,

- 10 (CR³R^{3a})₀₋₂CR³=CR³(CR³R^{3a})₀₋₂, (CR³R^{3a})₀₋₂C≡C(CR³R^{3a})₀₋₂,
(CR³R^{3a})_uC(O)(CR³R^{3a})_w, (CR³R^{3a})_uC(O)O(CR³R^{3a})_w,
(CR³R^{3a})_uOC(O)(CR³R^{3a})_w, (CR³R^{3a})_uO(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_w, (CR³R^{3a})_uOC(O)NR^{3b}(CR³R^{3a})_w,
15 (CR³R^{3a})_uNR^{3b}C(O)O(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(S)NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uS(CR³R^{3a})_w,
(CR³R^{3a})_uS(O)(CR³R^{3a})_w, (CR³R^{3a})_uS(O)₂(CR³R^{3a})_w,
(CR³R^{3a})_uS(O)NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uNR^{3b}S(O)₂(CR³R^{3a})_w,
20 (CR³R^{3a})_uS(O)₂NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}S(O)₂NR^{3b}(CR³R^{3a})_w, (CR³R^{3a})_uNR^{3e}(CR³R^{3a})_w,
(CR³R^{3a})_uC(O)(CR³R^{3a})_uC(O)(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(O)(CR³R^{3a})_w,
25 (CR³R^{3a})_uC(O)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3bb}C(S)(CR³R^{3a})_uC(O)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uNR^{3b}C(O)(CR³R^{3a})_uC(S)NR^{3b}(CR³R^{3a})_w,
(CR³R^{3a})_uS(O)NR^{3b}C(O)(CR³R^{3a})_w,
30 (CR³R^{3a})_uC(O)NR^{3b}S(O)₂(CR³R^{3a})_w, and
(CR³R^{3a})_uS(O)₂NR^{3b}C(O)NR^{3b}(CR³R^{3a})_w, wherein u + w total
0, 1, 2, 3, or 4, provided that G₁ does not form a N-

S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

Z is selected from a bond, $-(CR^3R^3e)_{1-4}-$,

- 5 $(CR^3R^3e)_qO(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qNR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qC(O)(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qC(O)O(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qOC(O)(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qC(O)NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}C(O)(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qOC(O)O(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qOC(O)NR^{3b}(CR^3R^3e)_{q1}$,
10 $(CR^3R^3e)_qNR^{3b}C(O)O(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}C(O)NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qC(O)(CR^3R^3e)_qC(O)(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}(CR^3R^3e)_qC(O)NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}C(O)(CR^3R^3e)_qC(O)(CR^3R^3e)_{q1}$,
15 $(CR^3R^3e)_qC(O)(CR^3R^3e)_qC(O)NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}C(O)(CR^3R^3e)_qC(O)NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qS(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qS(O)(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qS(O)_2(CR^3R^3e)_{q1}$, $(CR^3R^3e)_qSO_2NR^{3b}(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qNR^{3b}SO_2(CR^3R^3e)_{q1}$,
20 $(CR^3R^3e)_qS(O)NR^{3b}C(O)(CR^3R^3e)_{q1}$,
 $(CR^3R^3e)_qC(O)NR^{3b}S(O)_2(CR^3R^3e)_{q1}$, and
 $(CR^3R^3e)_qNR^{3b}SO_2NR^{3b}(CR^3R^3e)_{q1}$, wherein q + q1 total 0,
1, 2, 3, or 4, provided that Z does not form a N-S,
NCH₂N, NCH₂O, or NCH₂S bond with either group to which
25 it is attached;

provided that:

- (a) when ring P is absent and ring M is a pyridyl ring, then Z is other than C(O)NHCH₂; and,
30 (b) when ring P is absent and ring M is a piperazinyl ring, then either Z is other than alkylene or A is other than phenyl;

Z^2 is selected from H, $S(O)_2NHR^{3b}$, $C(O)R^{3b}$, $C(O)NHR^{3b}$,
 $C(O)OR^{3f}$, $S(O)R^{3f}$, $S(O)_2R^{3f}$, C_{1-6} alkyl substituted with
 0-2 R^{1a} , C_{2-6} alkenyl substituted with 0-2 R^{1a} , C_{2-6}
 alkynyl substituted with 0-2 R^{1a} , $-(C_{0-4} \text{ alkyl})-\dot{C}_{3-10}$
 carbocycle substituted with 0-3 R^{1a} , and $-(C_{0-4} \text{ alkyl})-$
 5 5-10 membered heterocycle substituted with 0-3 R^{1a} and
 consisting of: carbon atoms and 1-4 heteroatoms
 selected from the group consisting of N, O, and $S(O)_p$;

10 R^{1a} , at each occurrence, is selected from H, $-(CR^3R^{3a})_r-R^{1b}$,
 $-(CR^3R^{3a})_r-CR^3R^{1b}R^{1b}$, $-(CR^3R^{3a})_r-O-(CR^3R^{3a})_r-R^{1b}$,
 $-(CR^3R^{3a})_r-NR^2-(CR^3R^{3a})_r-R^{1b}$,
 $-(CR^3R^{3a})_r-S(O)_p-(CR^3R^{3a})_r-R^{1b}$,
 $-(CR^3R^{3a})_r-CO_2-(CR^3R^{3a})_r-R^{1b}$,
 15 $-(CR^3R^{3a})_r-C(O)NR^2-(CR^3R^{3a})_r-R^{1b}$,
 $-(CR^3R^{3a})_r-C(O)-(CR^3R^{3a})_r-R^{1b}$, $-C_{2-6}$ alkenylene- R^{1b} ,
 $-C_{2-6}$ alkynylene- R^{1b} , and $-(CR^3R^{3a})_r-C(=NR^{1b})NR^3R^{1b}$,
 provided that R^{1a} forms other than an N-halo, N-S, O-O,
 or N-CN bond;

20 alternatively, when two R^{1a} groups are attached to adjacent
 atoms or to the same carbon atom, together with the
 atoms to which they are attached, they form a 5-7
 membered ring consisting of: carbon atoms and 0-2
 25 heteroatoms selected from the group consisting of N,
 O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b}
 and comprising: 0-3 double bonds;

R^{1b} is selected from H, C_{1-3} alkyl, F, Cl, Br, I, $-CN$, $-NO_2$,
 30 $-CHO$, $(CF_2)_rCF_3$, $(CR^3R^{3a})_rOR^2$, NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} ,
 $OC(O)R^2$, $CH(CH_2OR^2)_2$, $(CF_2)_rCO_2R^{2a}$, $S(O)_pR^{2b}$,
 $NR^2(CH_2)_rOR^2$, $C(=NR^{2c})NR^2R^{2a}$, $NR^2C(O)R^{2b}$, $NR^2C(O)NR^2R^{2a}$,
 $NR^2C(O)_2R^{2a}$, $OC(O)NR^2R^{2a}$, $C(O)NR^2R^{2a}$, $C(O)NR^2(CH_2)_rOR^2$,

SO₂NR²R^{2a}, NR²SO₂R², C(O)NR²SO₂R², C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group
5 consisting of N, O, and S(O)_p, provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond and provided that S(O)_pR² forms other than S(O)₂H or S(O)H;

R², at each occurrence, is selected from H, CF₃, C₁₋₆ alkyl, benzyl, -(CH₂)_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and -(CH₂)_r-5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4
10 heteroatoms selected from the group consisting of N, O, and S(O)_p;

15 R^{2a}, at each occurrence, is selected from H, CF₃, C₁₋₆ alkyl, benzyl, -(CH₂)_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and -(CH₂)_r-5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon
20 atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 3-6
25 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

30 R^{2b}, at each occurrence, is selected from CF₃, C₁₋₄ alkoxy substituted with 0-2 R^{4b}, C₁₋₆ alkyl substituted with 0-3 R^{4b}, -(CH₂)_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and -(CH₂)_r-5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4

heteroatoms selected from the group consisting of N, O, and S(O)_p;

- 5 R^{2c}, at each occurrence, is selected from CF₃, OH, C₁₋₄ alkoxy, C₁₋₆ alkyl, -(CH₂)_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and -(CH₂)_r-5-10 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;
- 10 R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₆ alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4c}, and -(CR³R^{3a})_r-5-10 membered heterocycle substituted with 0-2 R^{4c} and consisting of:
- 15 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;
- 20 alternatively, when two R^{2d}'s are attached to the same nitrogen atom, then R^{2d} and R^{2d}, together with the nitrogen atom to which they are attached, combine to form a 5-10 membered saturated, partially saturated or
- 25 unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;
- 30 R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₆ alkyl substituted with 0-2 R^{4c}, -(CR³R^{3a})_r-C₃₋₁₀ carbocycle substituted with 0-2 R^{4c}, and -(CR³R^{3a})_r-5-10 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the

group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

5 R³, at each occurrence, is selected from H, CH₃, CH₂CH₃,
CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂,
CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, and phenyl;

10 R^{3a}, at each occurrence, is selected from H, CH₃, CH₂CH₃,
CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂,
CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, and phenyl;

alternatively, R³ and R^{3a}, together with the nitrogen atom
to which they are attached, combine to form a 5 or 6
membered saturated, partially unsaturated, or
15 unsaturated ring consisting of: carbon atoms, the
nitrogen atom to which R³ and R^{3a} are attached, and 0-1
additional heteroatoms selected from the group
consisting of N, O, and S(O)_p;

20 R^{3b}, at each occurrence, is selected from H, C₁₋₆ alkyl
substituted with 0-2 R^{1a}, C₂₋₆ alkenyl substituted with
0-2 R^{1a}, C₂₋₆ alkynyl substituted with 0-2 R^{1a}, -(C₀₋₄
alkyl)-5-10 membered carbocycle substituted with 0-3
R^{1a}, and -(C₀₋₄ alkyl)- 5-10 membered heterocycle
25 substituted with 0-3 R^{1a} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and S(O)_p;

30 R^{3c}, at each occurrence, is selected from CH₃, CH₂CH₃,
CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂,
CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, and phenyl;

R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, C_{1-4} alkyl-phenyl, and $C(=O)R^{3c}$;

5 R^{3e} , at each occurrence, is selected from H, $S(O)_2NHR^3$, $C(O)R^3$, $C(O)NHR^3$, $C(O)OR^{3f}$, $S(O)R^{3f}$, $S(O)_2R^{3f}$, C_{1-6} alkyl substituted with 0-2 R^{1a} , C_{2-6} alkenyl substituted with 0-2 R^{1a} , C_{2-6} alkynyl substituted with 0-2 R^{1a} , $-(C_{0-4}$ alkyl)-5-10 membered carbocycle substituted with 0-3 R^{1a} , and $-(C_{0-4}$ alkyl)-5-10 membered heterocycle substituted with 0-3 R^{1a} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

15 R^{3f} , at each occurrence, is selected from: C_{1-6} alkyl substituted with 0-2 R^{1a} , C_{2-6} alkenyl substituted with 0-2 R^{1a} , C_{2-6} alkynyl substituted with 0-2 R^{1a} , $-(C_{0-4}$ alkyl)-5-10 membered carbocycle substituted with 0-3 R^{1a} , and $-(C_{0-4}$ alkyl)-5-10 membered heterocycle substituted with 0-3 R^{1a} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

25 R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, $-(CH_2)_r$ -3-6 membered carbocycle, and $-(CH_2)_r$ -5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

30

alternatively, when R^3 and R^{3g} are attached to the same carbon atom, they combine with the attached carbon atom to form a cyclopropyl group;

R^4 , at each occurrence, is selected from H, =O,
 $(CR^3R^{3a})_rOR^2$, F, Cl, Br, I, C_{1-4} alkyl, $(CR^3R^{3a})_rCN$,
 $(CR^3R^{3a})_rNO_2$, $(CR^3R^{3a})_rNR^2R^{2a}$, $(CR^3R^{3a})_rC(O)R^{2c}$,
5 $(CR^3R^{3a})_rNR^2C(O)R^{2b}$, $(CR^3R^{3a})_rC(O)NR^2R^{2a}$,
 $(CR^3R^{3a})_rNR^2C(O)NR^2R^{2a}$, $(CR^3R^{3a})_rC(=NR^2)NR^2R^{2a}$,
 $(CR^3R^{3a})_rC(=NS(O)_2R^{5a})NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2C(=NR^2)NR^2R^{2a}$,
 $(CR^3R^{3a})_rC(O)NR^2C(=NR^2)NR^2R^{2a}$, $(CR^3R^{3a})_rSO_2NR^2R^{2a}$,
 $(CR^3R^{3a})_rNR^2SO_2NR^2R^{2a}$, $(CR^3R^{3a})_rNR^2SO_2-C_{1-4}$ alkyl,
10 $(CR^3R^{3a})_rNR^2SO_2R^{5a}$, $(CR^3R^{3a})_rS(O)_pR^{5a}$, $(CR^3R^{3a})_r(CF_2)_rCF_3$,
 $N(CH_2)_rR^{1b}$, $O(CH_2)_rR^{1b}$, $S(CH_2)_rR^{1b}$, $(CR^3R^{3a})_{r-5-6}$
membered carbocycle substituted with 0-1 R^5 , and a
 $(CR^3R^{3a})_{r-5-6}$ membered heterocycle substituted with 0-1
 R^5 and consisting of: carbon atoms and 1-4
15 heteroatoms selected from the group consisting of N,
O, and $S(O)_p$;

R^{4a} is selected from C_{1-6} alkyl substituted with 0-2 R^{4c} ,
 C_{2-6} alkenyl substituted with 0-2 R^{4c} , C_{2-6} alkynyl
20 substituted with 0-2 R^{4c} , $-(CR^3R^{3g})_r-C_{5-10}$ membered
carbocycle substituted with 0-3 R^{4c} , $-(CR^3R^{3g})_{r-5-10}$
membered heterocycle substituted with 0-3 R^{4c} and
consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and $S(O)_p$,
25 $(CR^3R^{3g})_rCN$, $(CR^3R^{3g})_rC(=NR^{2d})NR^{2d}R^{2d}$,
 $(CR^3R^{3g})_rNR^{2d}C(=NR^{2d})NR^{2d}R^{2d}$, $(CR^3R^{3g})_rNR^{2d}C(R^{2e})(=NR^{2d})$,
 $(CR^3R^{3g})_rNR^{2d}R^{2d}$, $(CR^3R^{3g})_rN(\rightarrow O)R^{2d}R^{2d}$, $(CR^3R^{3g})_rOR^{2d}$,
 $(CR^3R^{3g})_r-NR^{2d}C(O)R^{2e}$, $(CR^3R^{3g})_r-C(O)R^{2e}$,
 $(CR^3R^{3g})_r-OC(O)R^{2e}$, $(CR^3R^{3g})_r-C(O)NR^{2d}R^{2d}$,
30 $(CR^3R^{3g})_r-C(O)OR^{2d}$, $(CR^3R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d}$,
 $(CR^3R^{3g})_r-OC(O)NR^{2d}R^{2d}$, $(CR^3R^{3g})_r-NR^{2d}C(O)OR^{2d}$,
 $(CR^3R^{3g})_r-SO_2NR^{2d}R^{2d}$, $(CR^3R^{3g})_r-NR^{2d}SO_2NR^{2d}R^{2d}$,

$(\text{CR}^3\text{R}^3\text{g})_r\text{-C(O)NR}^{2d}\text{SO}_2\text{R}^{2d}$, $(\text{CR}^3\text{R}^3\text{g})_r\text{-NR}^{2d}\text{SO}_2\text{R}^{2d}$, and $(\text{CR}^3\text{R}^3\text{g})_r\text{-S(O)}_p\text{R}^{2d}$, provided that $\text{S(O)}_p\text{R}^{2d}$ forms other than $\text{S(O)}_2\text{H}$ or S(O)H and further provided that R^{4a} is other than a hydroxamic acid;

5

R^{4b} , at each occurrence, is selected from H, =O, $(\text{CH}_2)_r\text{OR}^3$, $(\text{CH}_2)_r\text{F}$, $(\text{CH}_2)_r\text{Cl}$, $(\text{CH}_2)_r\text{Br}$, $(\text{CH}_2)_r\text{I}$, C₁₋₄ alkyl, $(\text{CH}_2)_r\text{CN}$, $(\text{CH}_2)_r\text{NO}_2$, $(\text{CH}_2)_r\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{C(O)R}^3$, $(\text{CH}_2)_r\text{C(O)OR}^{3c}$, $(\text{CH}_2)_r\text{NR}^3\text{C(O)R}^{3a}$, $(\text{CH}_2)_r\text{-C(O)NR}^3\text{R}^{3a}$,
 10 $(\text{CH}_2)_r\text{NR}^3\text{C(O)NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{-C(=NR}^3)\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{C(=NR}^3)\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{SO}_2\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{NR}^3\text{R}^{3a}$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{-C}_{1-4}\text{ alkyl}$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{CF}_3$, $(\text{CH}_2)_r\text{NR}^3\text{SO}_2\text{-phenyl}$, $(\text{CH}_2)_r\text{S(O)}_p\text{CF}_3$, $(\text{CH}_2)_r\text{S(O)}_p\text{-C}_{1-4}\text{ alkyl}$, $(\text{CH}_2)_r\text{S(O)}_p\text{-phenyl}$, and
 15 $(\text{CH}_2)_r(\text{CF}_2)_r\text{CF}_3$;

R^{4c} , at each occurrence, is selected from =O, $(\text{CR}^3\text{R}^{3a})_r\text{OR}^2$, $(\text{CR}^3\text{R}^{3a})_r\text{F}$, $(\text{CR}^3\text{R}^{3a})_r\text{Br}$, $(\text{CR}^3\text{R}^{3a})_r\text{Cl}$, $(\text{CR}^3\text{R}^{3a})_r\text{CF}_3$, C₁₋₄ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, $(\text{CR}^3\text{R}^{3a})_r\text{CN}$,
 20 $(\text{CR}^3\text{R}^{3a})_r\text{NO}_2$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{N(}\rightarrow\text{O)R}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{C(O)R}^{2c}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C(O)R}^{2b}$, $(\text{CR}^3\text{R}^{3a})_r\text{C(O)NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{N=CHOR}^3$, $(\text{CR}^3\text{R}^{3a})_r\text{C(O)NR}^2(\text{CH}_2)_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C(O)NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{C(=NR}^2)\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{C(=NR}^2)\text{NR}^2\text{R}^{2a}$,
 25 $(\text{CR}^3\text{R}^{3a})_r\text{SO}_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})_r\text{C(O)NR}^2\text{SO}_2\text{-C}_{1-4}\text{ alkyl}$, $(\text{CR}^3\text{R}^{3a})_r\text{NR}^2\text{SO}_2\text{R}^{5a}$, $(\text{CR}^3\text{R}^{3a})_r\text{S(O)}_p\text{R}^{5a}$, $(\text{CF}_2)_r\text{CF}_3$, $(\text{CR}^3\text{R}^{3a})_r\text{C}_{3-10}\text{ carbocycle}$ substituted with 0-2 R^{4b} , and $(\text{CR}^3\text{R}^{3a})_r\text{4-10 membered}$ heterocycle substituted with 0-2 R^{4b} and consisting of
 30 carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p ;

R^5 , at each occurrence, is selected from H, C_{1-6} alkyl, =O, $(CH_2)_rOR^3$, F, Cl, Br, I, -CN, NO_2 , $(CH_2)_rNR^3R^{3a}$, $(CH_2)_rC(O)R^3$, $(CH_2)_rC(O)OR^{3c}$, $(CH_2)_rNR^3C(O)R^{3a}$, $(CH_2)_rC(O)NR^3R^{3a}$, $(CH_2)_rNR^3C(O)NR^3R^{3a}$, $(CH_2)_rCH(=NOR^{3d})$,
 5 $(CH_2)_rC(=NR^3)NR^3R^{3a}$, $(CH_2)_rNR^3C(=NR^3)NR^3R^{3a}$, $(CH_2)_rSO_2NR^3R^{3a}$, $(CH_2)_rNR^3SO_2NR^3R^{3a}$, $(CH_2)_rNR^3SO_2-C_{1-4}$ alkyl, $(CH_2)_rNR^3SO_2CF_3$, $(CH_2)_rNR^3SO_2$ -phenyl, $(CH_2)_rS(O)_pCF_3$, $(CH_2)_rS(O)_p-C_{1-4}$ alkyl, $(CH_2)_rS(O)_p$ -phenyl, $(CF_2)_rCF_3$, phenyl substituted with 0-2 R^6 ,
 10 naphthyl substituted with 0-2 R^6 , and benzyl substituted with 0-2 R^6 ;

R^{5a} , at each occurrence, is selected from C_{1-6} alkyl, $(CH_2)_rOR^3$, $(CH_2)_rNR^3R^{3a}$, $(CH_2)_rC(O)R^3$, $(CH_2)_rC(O)OR^{3c}$,
 15 $(CH_2)_rNR^3C(O)R^{3a}$, $(CH_2)_rC(O)NR^3R^{3a}$, $(CF_2)_rCF_3$, phenyl substituted with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and benzyl substituted with 0-2 R^6 , provided that R^{5a} does not form a S-N or $S(O)_p-C(O)$ bond;

20 R^6 , at each occurrence, is selected from H, OH, $(CH_2)_rOR^2$, halo, C_{1-4} alkyl, -CN, NO_2 , $(CH_2)_rNR^2R^{2a}$, $(CH_2)_rC(O)R^{2b}$, $NR^2C(O)R^{2b}$, $NR^2C(O)NR^2R^{2a}$, $C(=NH)NH_2$, $NHC(=NH)NH_2$, $SO_2NR^2R^{2a}$, $NR^2SO_2NR^2R^{2a}$, and $NR^2SO_2C_{1-4}$ alkyl;

25 R^7 , at each occurrence, is selected from H, OH, C_{1-6} alkyl, C_{1-6} alkyl-C(O)-, C_{1-6} alkyl-O-, $(CH_2)_n$ -phenyl, C_{1-4} alkyl-OC(O)-, C_{6-10} aryl-O-, C_{6-10} aryl-OC(O)-, C_{6-10} aryl-CH₂-C(O)-, C_{1-4} alkyl-C(O)O- C_{1-4} alkyl-OC(O)-, C_{6-10} aryl-C(O)O- C_{1-4} alkyl-OC(O)-,
 30 C_{1-6} alkyl-NH₂-C(O)-, phenyl-NH₂-C(O)-, and phenyl C_{1-4} alkyl-C(O)-;

R⁸, at each occurrence, is selected from H, C₁₋₆ alkyl, and (CH₂)_n-phenyl;

alternatively, R⁷ and R⁸, when attached to the same
 5 nitrogen, combine to form a 5-10 membered heterocyclic ring consisting of carbon atoms and 0-2 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

10 R⁹, at each occurrence, is selected from H, C₁₋₆ alkyl, and (CH₂)_n-phenyl;

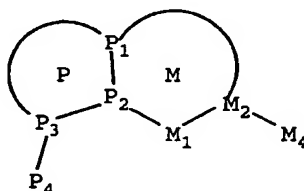
n, at each occurrence, is selected from 0, 1, 2, and 3;

15 p, at each occurrence, is selected from 0, 1, and 2;

r, at each occurrence, is selected from 0, 1, 2, 3, 4, 5, and 6; and,

20 t, at each occurrence, is selected from 0, 1, 2, and 3.

2. A compound according to Claim 1, wherein the compound is of Formula II:



25

II

or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

30 ring M, including P₁, P₂, M₁, and M₂, is a 5, 6, or 7 membered carbocycle or a 5, 6, or 7 membered

heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, N, and NZ²;

5 ring M is substituted with 0-2 R^{1a} and 0-2 carbonyl groups, and there are 0-3 ring double bonds;

ring P, including P₁, P₂, and P₃, is a 5 or 6 membered aromatic heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

10

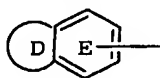
alternatively, ring P, including P₁, P₂, and P₃, is a 5 or 6 membered dihydro-aromatic heterocycle, consisting of: carbon atoms and 1-3 heteroatoms selected from O, S(O)_p, and N;

15

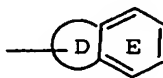
ring P is substituted with 0-2 R^{1a};

one of P₄ and M₄ is -Z-A-B and the other -G₁-G;

20 G is a group of formula IIa or IIb:



IIa



IIb

25 ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;

30

E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

alternatively, ring D is absent, and ring E is selected from phenyl, pyridyl, pyrimidyl, and thienyl, and ring E is substituted with 1-3 R;

- 5 alternatively, ring D is absent, ring E is selected from phenyl, pyridyl, and thienyl, and ring E is substituted with 1 R and substituted with a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of
 10 N, O, and S(O)_p, wherein the 5-6 membered heterocycle is substituted with 0-2 carbonyls and 1-3 R and there are 0-3 ring double bonds;

- R is selected from H, C₁₋₄ alkyl, F, Cl, OH, OCH₃, OCH₂CH₃,
 15 OCH(CH₃)₂, CN, C(=NH)NH₂, C(=NH)NHOH, C(=NH)NHOCH₃, NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂, CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tNR⁷R⁸, C(O)NR⁷R⁸, CH₂C(O)NR⁷R⁸, S(O)_pNR⁷R⁸, CH₂S(O)_pNR⁷R⁸, SO₂R³, and OCF₃;

- 20 alternatively, when 2 R groups are attached to adjacent atoms, they combine to form methylenedioxy or ethylenedioxy;

- 25 A is selected from:
 C₅₋₁₀ carbocycle substituted with 0-2 R⁴, and
 5-10 membered heterocycle substituted with 0-2 R⁴ and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

- 30 X is selected from -(CR²R^{2a})₁₋₄-, -C(O)-, -C(O)CR²R^{2a}-, -CR²R^{2a}C(O)-, -S(O)₂-, -S(O)₂CR²R^{2a}-, -CR²R^{2a}S(O)₂-, -NR²S(O)₂-, -S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR², -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

- Y is a C₃₋₇ monocyclic carbocycle or 3-7 membered monocyclic heterocycle, wherein the carbocycle or heterocycle consists of: carbon atoms and 0-2 heteroatoms selected from N, O, and S(O)_p, the carbocycle or heterocycle further comprises 0-2 double bonds and 0-2 carbonyl groups, and the carbocycle or heterocycle is substituted with 0-2 R⁴;
- alternatively, Y is CY¹Y², and Y¹ and Y² are independently C₁₋₃ alkyl substituted with 0-1 R⁴;
- Z is selected from a bond, CH₂, CH₂CH₂, CH₂O, OCH₂, C(O), NH, CH₂NH, NHCH₂, CH₂C(O), C(O)CH₂, C(O)NH, NHC(O), NHC(O)CH₂C(O)NH, S(O)₂, CH₂S(O)₂, S(O)₂(CH₂), SO₂NH, and NHSO₂, provided that Z does not form a N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;
- Z² is selected from H, C₁₋₄ alkyl, phenyl, benzyl, C(O)R^{3b}, S(O)R^{3f}, and S(O)₂R^{3f};
- R^{1a}, at each occurrence, is selected from H, -(CH₂)_r-R^{1b}, -(CH(CH₃))_r-R^{1b}, -(C(CH₃)₂)_r-R^{1b}, -O-(CR³R^{3a})_r-R^{1b}, -NR²-(CR³R^{3a})_r-R^{1b}, and -S-(CR³R^{3a})_r-R^{1b}, provided that R^{1a} forms other than an N-halo, N-S, O-O, or N-CN bond;
- alternatively, when two R^{1a} groups are attached to adjacent atoms or to the same carbon atom, together with the atoms to which they are attached, they form a 5-7 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p, this ring being substituted with 0-2 R^{4b} and comprising: 0-3 double ring bonds;

R^{1b} is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, F, Cl, Br, I, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , $S(O)_pR^{2b}$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$,
 5 $NR^2C(O)NHR^2$, $NR^2C(O)_2R^{2a}$, $OC(O)NR^2R^{2a}$, $C(O)NR^2R^{2a}$, $C(O)NR^2(CH_2)_rOR^2$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group
 10 consisting of N, O, and $S(O)_p$, provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond and provided that $S(O)_pR^2$ forms other than $S(O)_2H$ or $S(O)H$;

R^2 , at each occurrence, is selected from H, CF_3 , CH_3 ,
 15 CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , a C_{5-6} carbocycle- CH_2 -substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon
 20 atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 ,
 CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 25 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

30

alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated

ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

5 R^{2b} , at each occurrence, is selected from CF_3 , C_{1-4} alkoxy, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of:
10 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2c} , at each occurrence, is selected from CF_3 , OH, C_{1-4} alkoxy, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$,
15 $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

20 R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of:
25 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p-S(O)_p$, S-O, O-N, O-S, or O-O moiety;

30 alternatively, when two R^{2d} 's are attached to the same nitrogen atom, then R^{2d} and R^{2d} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated

or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

- 5 R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the
10 group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a $C(O)$ -halo or $C(O)-S(O)_p$ moiety;

R^3 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

15

R^{3a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

- alternatively, R^3 and R^{3a} , together with the nitrogen atom
20 to which they are attached, combine to form a 5 or 6 membered saturated, partially unsaturated, or unsaturated ring consisting of: carbon atoms and the nitrogen atom to which R^3 and R^{3a} are attached;

- 25 R^{3c} , at each occurrence, is selected from CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

- R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, CH_2 -phenyl, CH_2CH_2 -phenyl, and
30 $C(=O)R^{3c}$;

R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, cyclopropyl-methyl, benzyl, and phenyl;

- 5 alternatively, when R^3 and R^{3g} are attached to the same carbon atom, they combine with the attached carbon atom to form a cyclopropyl group;

10 R^4 , at each occurrence, is selected from H, =O, OR^2 , CH_2OR^2 , $(CH_2)_2OR^2$, F, Cl, Br, I, C₁₋₄ alkyl, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $(CH_2)_2NR^2R^{2a}$, $C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , 5-6 membered carbocycle substituted with 0-1 R^5 , and a 5-6 membered heterocycle substituted with 0-1 R^5 and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{4b} , at each occurrence, is selected from H, =O, OR^3 , CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$,
20 $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, -CN, NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2-C(O)R^3$, $C(O)OR^{3c}$, $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $CH_2NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $CH_2C(O)NR^3R^{3a}$, $NR^3C(O)NR^3R^{3a}$, $CH_2NR^3C(O)NR^3R^{3a}$, $C(=NR^3)NR^3R^{3a}$, $CH_2C(=NR^3)NR^3R^{3a}$, $NR^3C(=NR^3)NR^3R^{3a}$,
25 $CH_2NR^3C(=NR^3)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $CH_2SO_2NR^3R^{3a}$, $NR^3SO_2NR^3R^{3a}$, $CH_2NR^3SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, $CH_2NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, $CH_2NR^3SO_2CF_3$, NR^3SO_2 -phenyl, $CH_2NR^3SO_2$ -phenyl, $S(O)_pCF_3$, $CH_2S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $CH_2S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl,
30 $CH_2S(O)_p$ -phenyl, CF_3 , and CH_2-CF_3 ;

R^{4c} , at each occurrence, is selected from =O, $(CR^3R^{3a})_rOR^2$, $(CR^3R^{3a})_rF$, $(CR^3R^{3a})_rBr$, $(CR^3R^{3a})_rCl$, $(CR^3R^{3a})_rCF_3$, C₁₋₄

alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, (CR³R^{3a})_rCN,
 (CR³R^{3a})_rNO₂, (CR³R^{3a})_rNR²R^{2a}, (CR³R^{3a})_rN(→O)R²R^{2a},
 (CR³R^{3a})_rC(O)R^{2c}, (CR³R^{3a})_rNR²C(O)R^{2b},
 (CR³R^{3a})_rC(O)NR²R^{2a}, (CR³R^{3a})_rNR²C(O)NR²R^{2a},
 5 (CR³R^{3a})_rSO₂NR²R^{2a}, (CR³R^{3a})_rNR²SO₂NR²R^{2a},
 (CR³R^{3a})_rNR²SO₂R^{5a}, (CR³R^{3a})_rS(O)_pR^{5a}, (CF₂)_rCF₃,
 (CR³R^{3a})_rC₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and
 (CR³R^{3a})_r5-10 membered heterocycle substituted with 0-2
 10 R^{4b} and consisting of carbon atoms and from 1-4
 heteroatoms selected from the group consisting of N,
 O, and S(O)_p;

R⁵, at each occurrence, is selected from H, =O, CH₃, CH₂CH₃,
 CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂,
 15 CH(CH₃)CH₂CH₃, C(CH₃)₃, OR³, CH₂OR³, F, Cl, -CN, NO₂,
 NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, CH₂C(O)R³, C(O)OR^{3c},
 CH₂C(O)OR^{3c}, NR³C(O)R^{3a}, C(O)NR³R^{3a}, NR³C(O)NR³R^{3a},
 CH(=NOR^{3d}), C(=NR³)NR³R^{3a}, NR³C(=NR³)NR³R^{3a}, SO₂NR³R^{3a},
 NR³SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, NR³SO₂CF₃, NR³SO₂-
 20 phenyl, S(O)_pCF₃, S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl, CF₃,
 phenyl substituted with 0-2 R⁶, naphthyl substituted
 with 0-2 R⁶, and benzyl substituted with 0-2 R⁶;

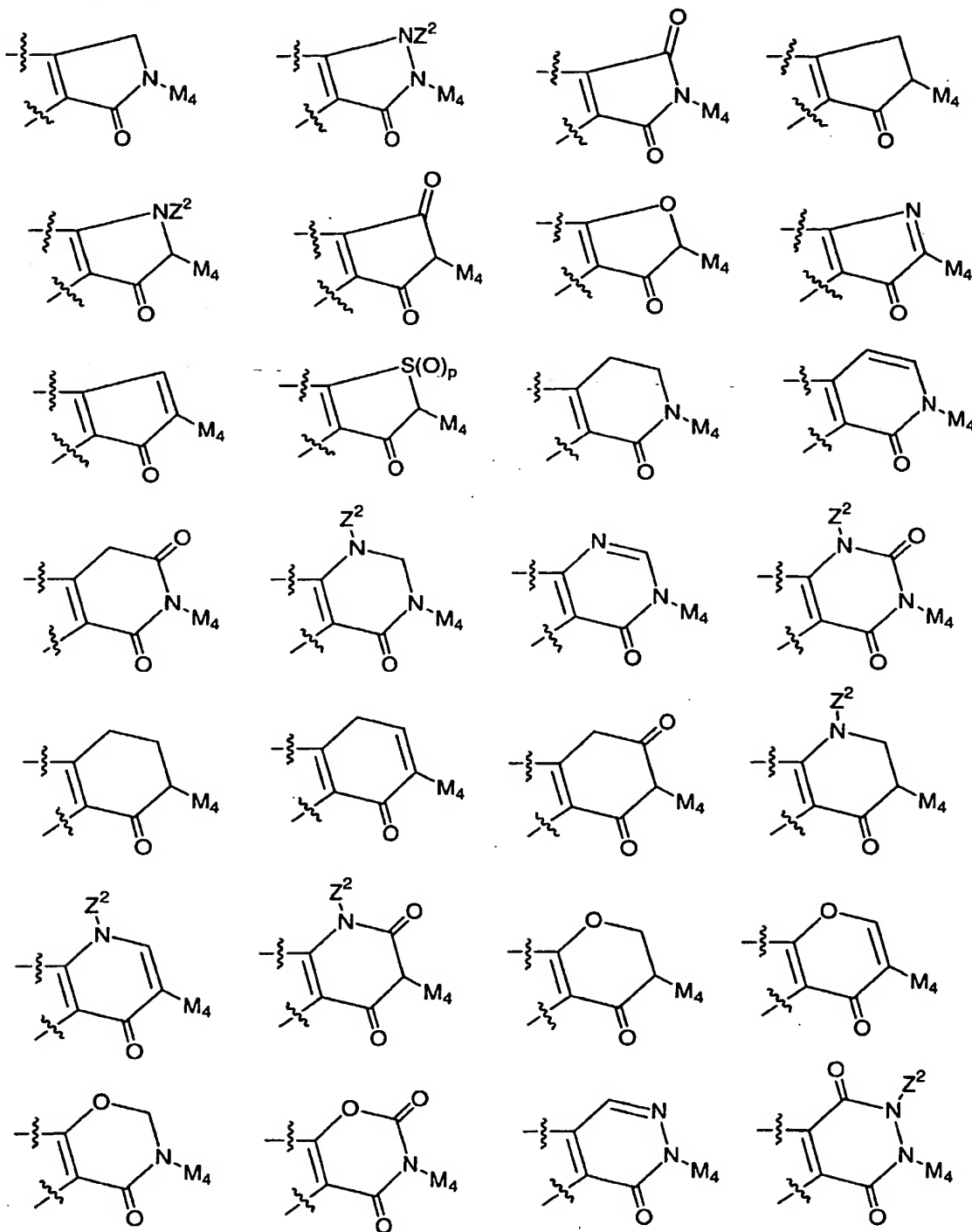
R⁶, at each occurrence, is selected from H, OH, OR², F, Cl,
 25 CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃,
 CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, -CN, NO₂, NR²R^{2a},
 CH₂NR²R^{2a}, C(O)R^{2b}, CH₂C(O)R^{2b}, NR²C(O)R^{2b},
 NR²C(O)NR²R^{2a}, C(=NH)NH₂, NHC(=NH)NH₂, SO₂NR²R^{2a},
 NR²SO₂NR²R^{2a}, and NR²SO₂-C₁₋₄ alkyl; and,

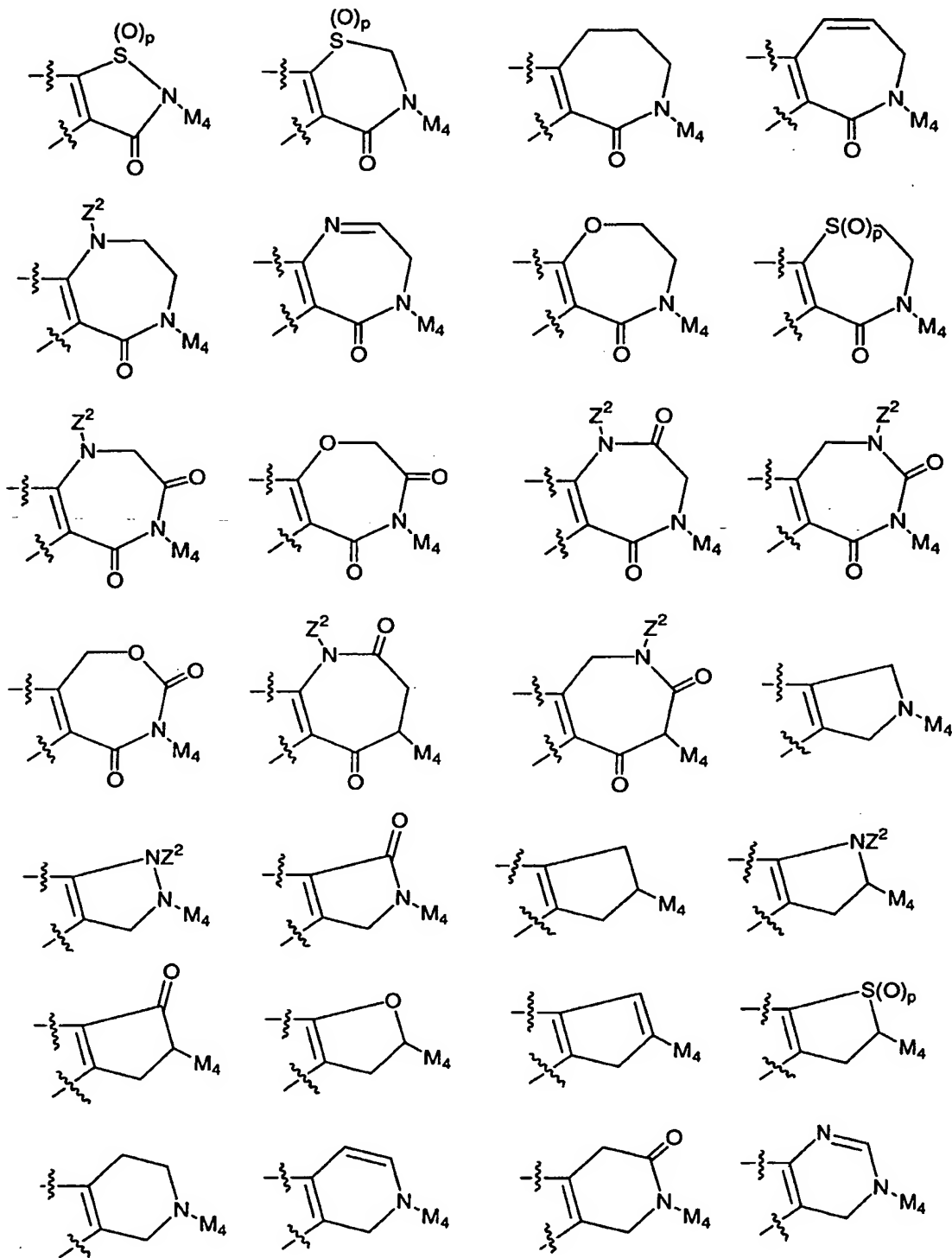
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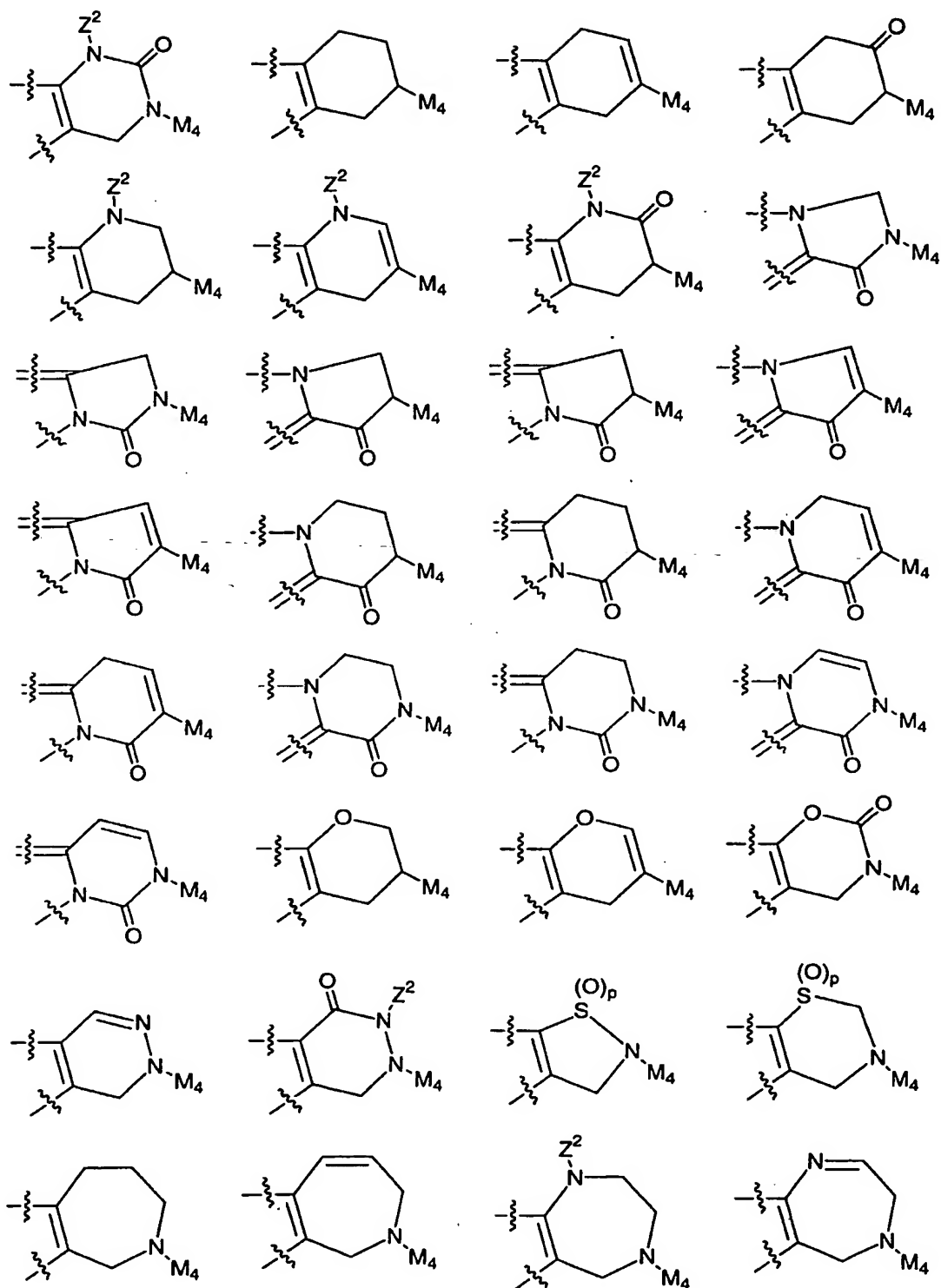
r, at each occurrence, is selected from 0, 1, 2, and 3.

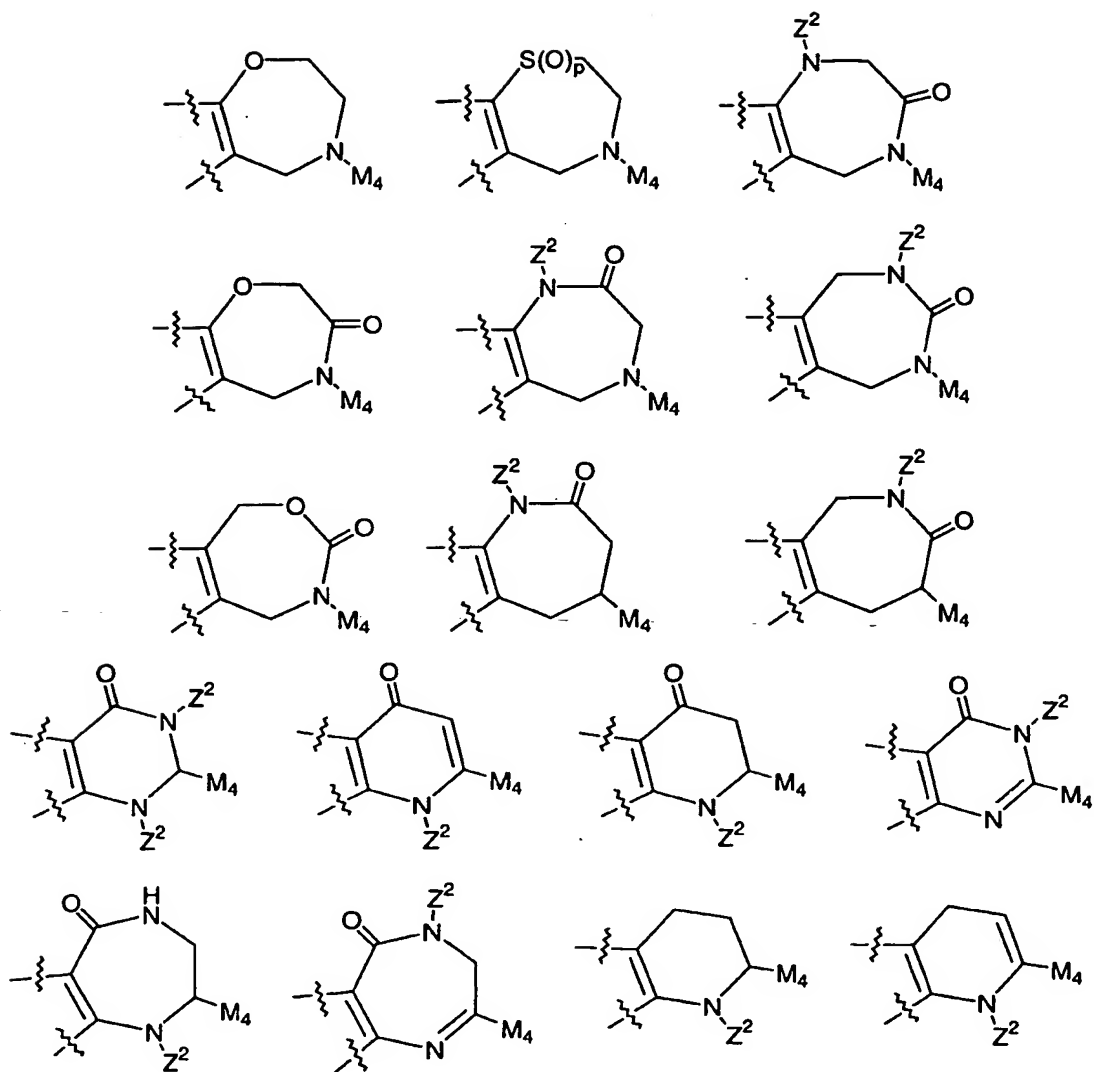
3. A compound according to Claim 2, wherein:

ring M is substituted with 0-2 R^{1a} and is selected from the group:

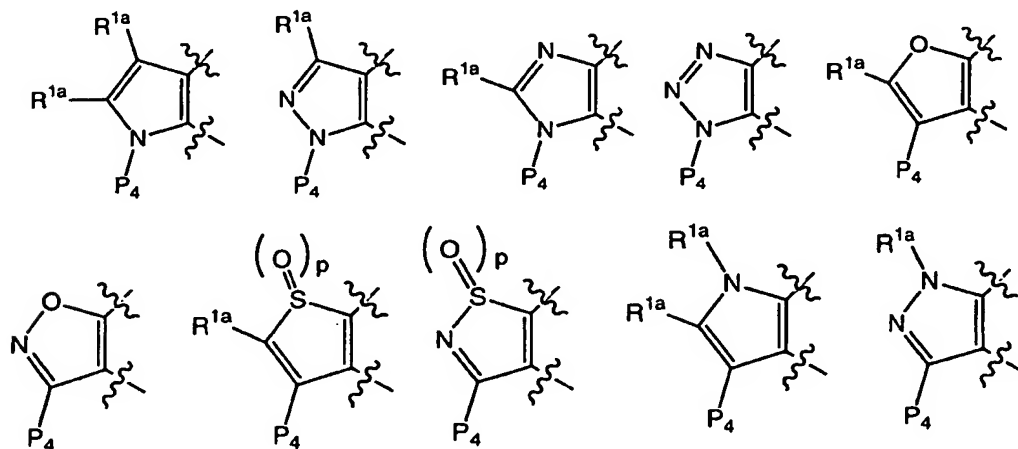


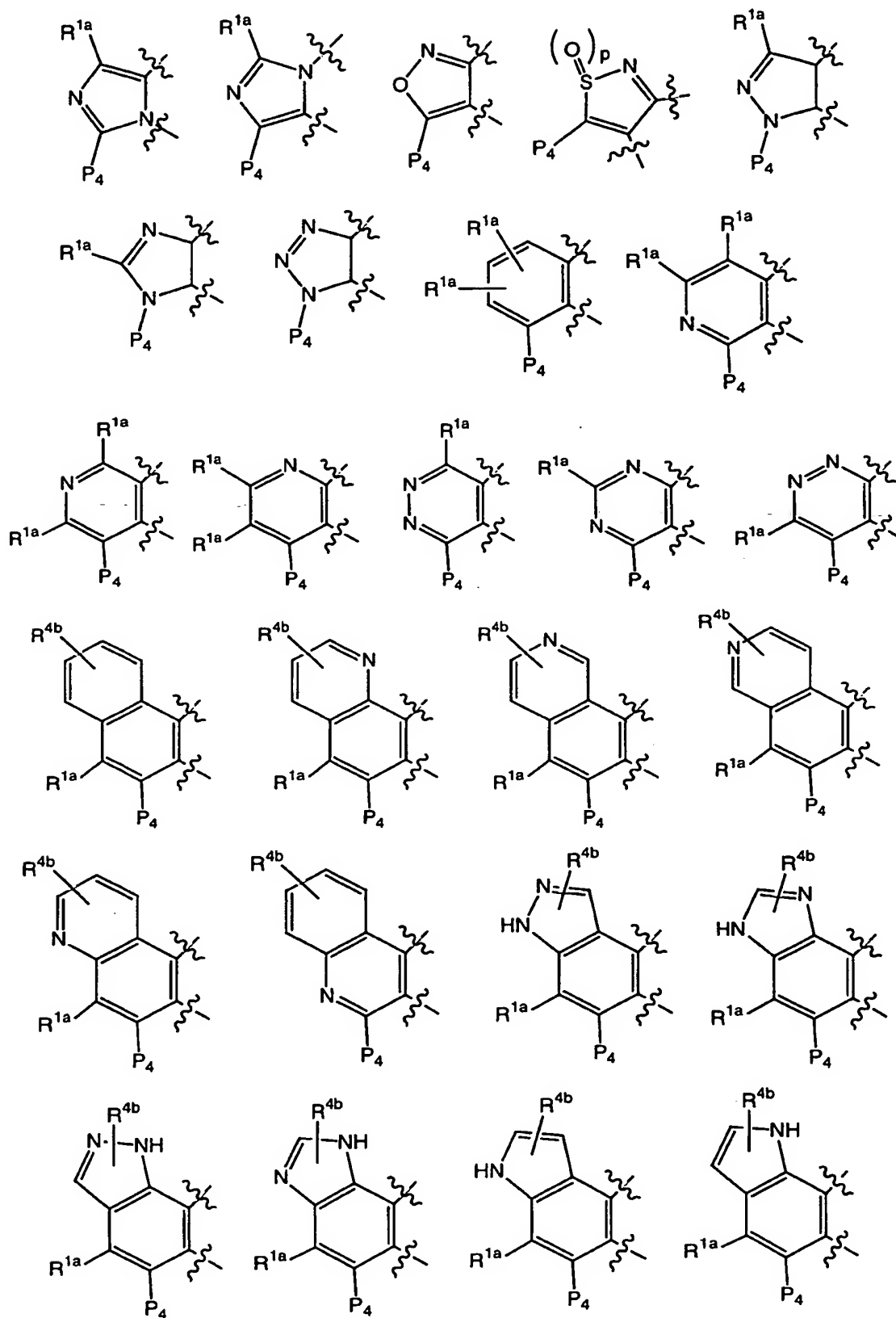


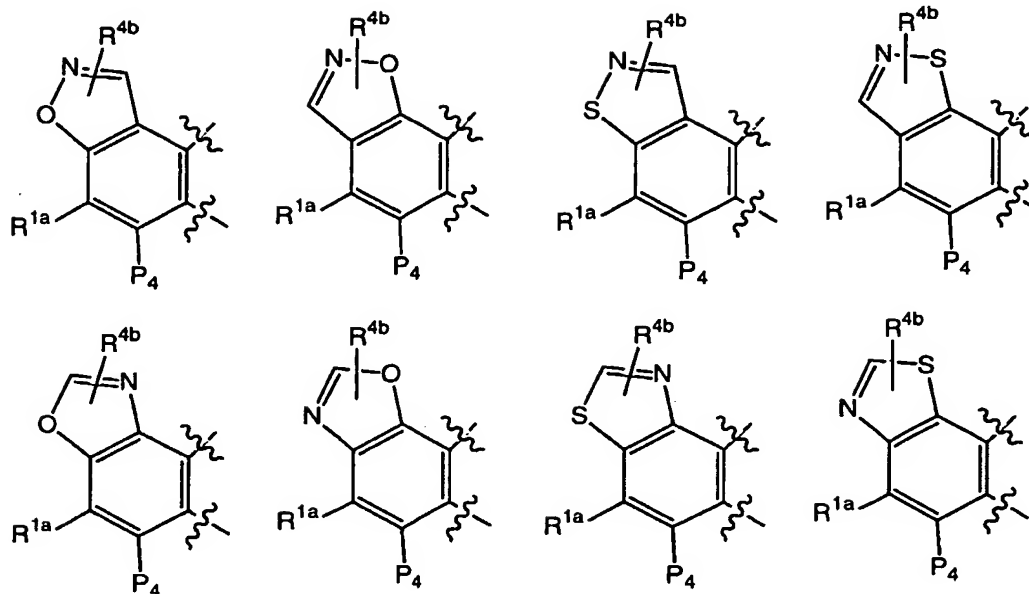




ring P, including P₁, P₂, P₃, and P₄ is selected from group:

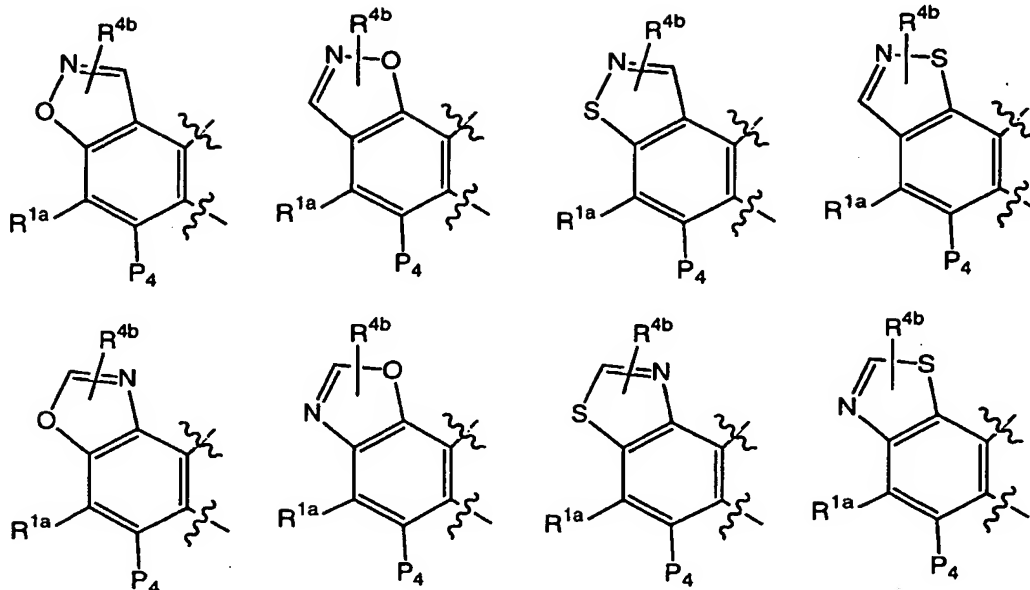






G is selected from the group:

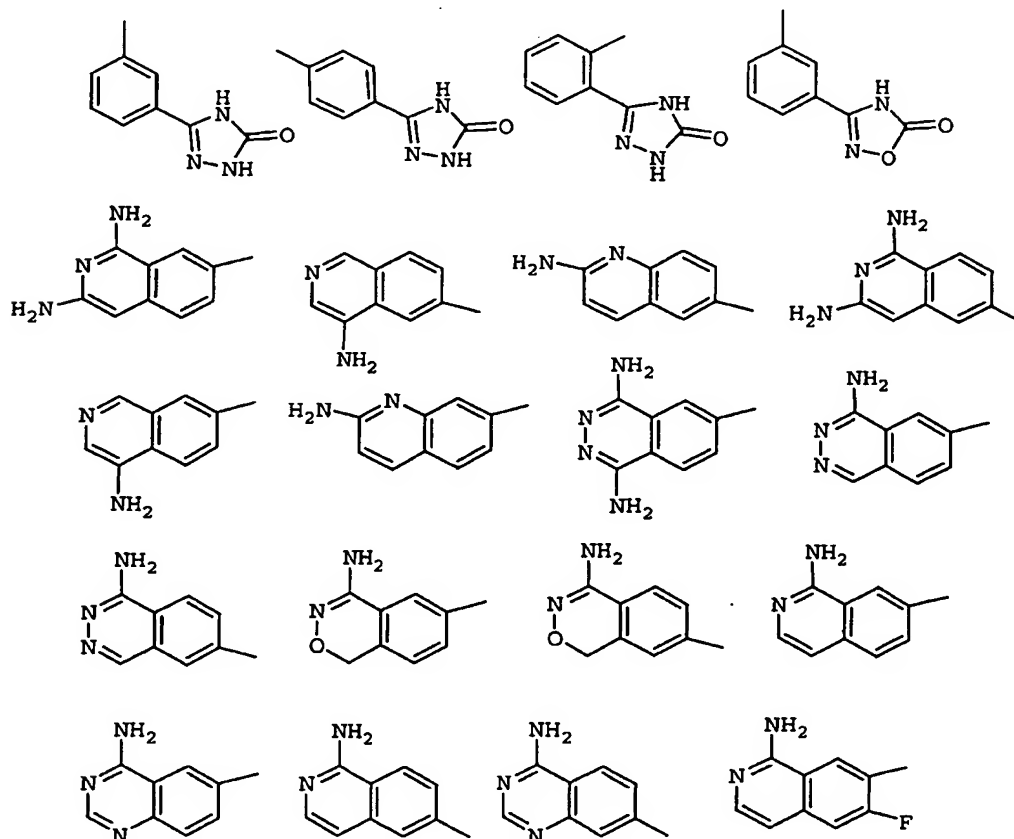
- phenyl; 2,5-bis-aminomethyl-phenyl;
- 5 2-amido-4-methoxy-phenyl; 2-amido-5-chloro-phenyl;
- 2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
- 2-aminomethyl-3-methoxy-phenyl;
- 2-aminomethyl-4-fluoro-phenyl;
- 2-aminomethyl-4-methoxy-phenyl;
- 10 2-aminomethyl-5-fluoro-phenyl;
- 2-aminomethyl-5-methoxy-phenyl;
- 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
- 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
- 2-aminosulfonyl-phenyl; 2-aminomethyl-4-ethyl-phenyl; 2-
- 15 aminosulfonyl-4-ethyl-phenyl; 2-amido-4-ethyl-phenyl;
- 2-hydroxy-4-methoxy-phenyl; 2-methylsulfonyl-phenyl;
- 3-(N,N-dimethylamino)-4-chloro-phenyl;
- 3-(N,N-dimethylamino)-phenyl; 3-(N-hydroxy-amidino)-phenyl;
- 3-(N-methoxy-amidino)-phenyl;
- 20 3-(N-methylamino)-4-chloro-phenyl;
- 3-(N-methylamino)-phenyl; 3-amidino-phenyl;
- 3-amido-6-hydroxy-phenyl; 3-amido-phenyl;
- 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
- 3-amino-phenyl; 3-chloro-4-fluoro-phenyl; 3-chloro-phenyl;

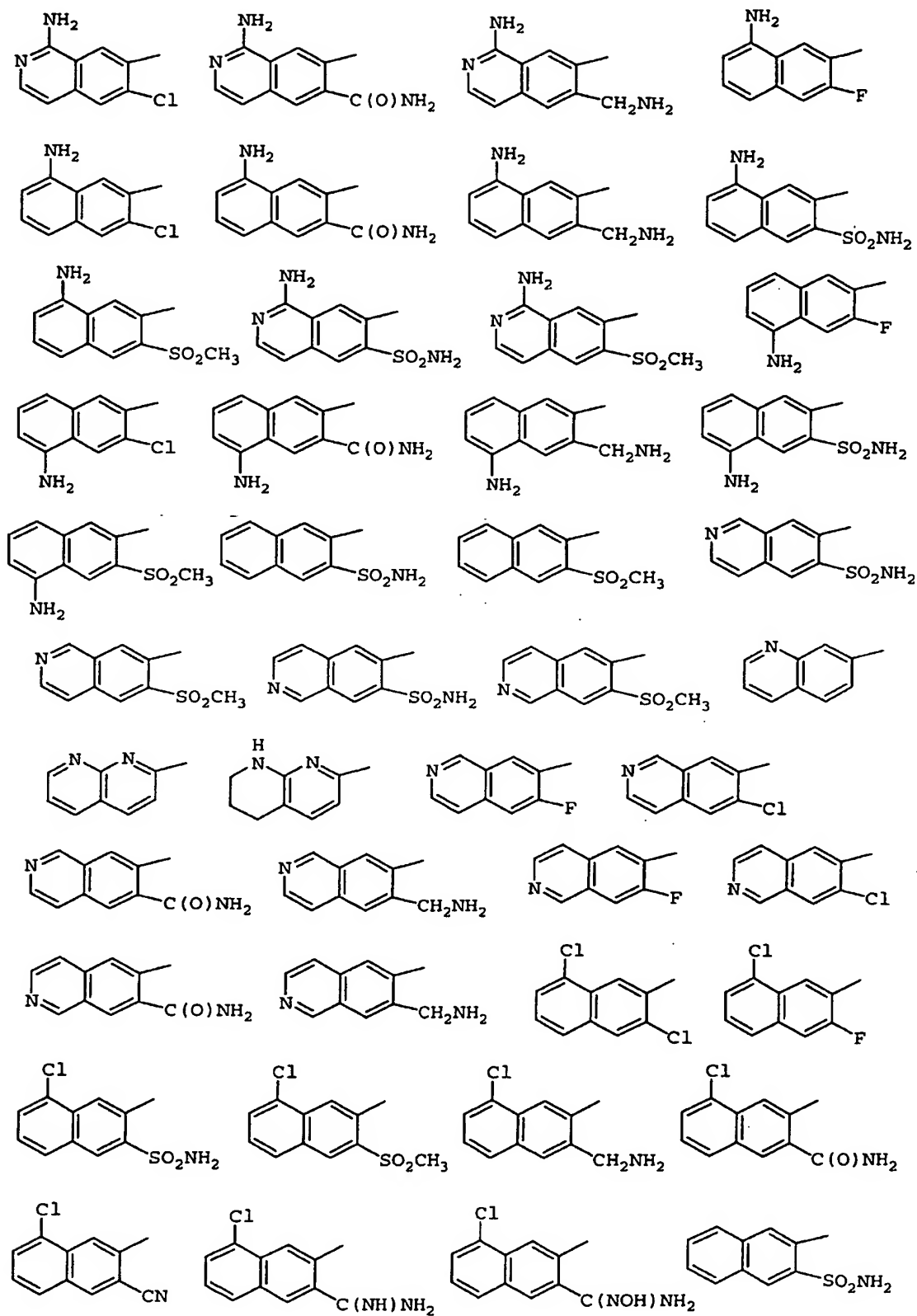


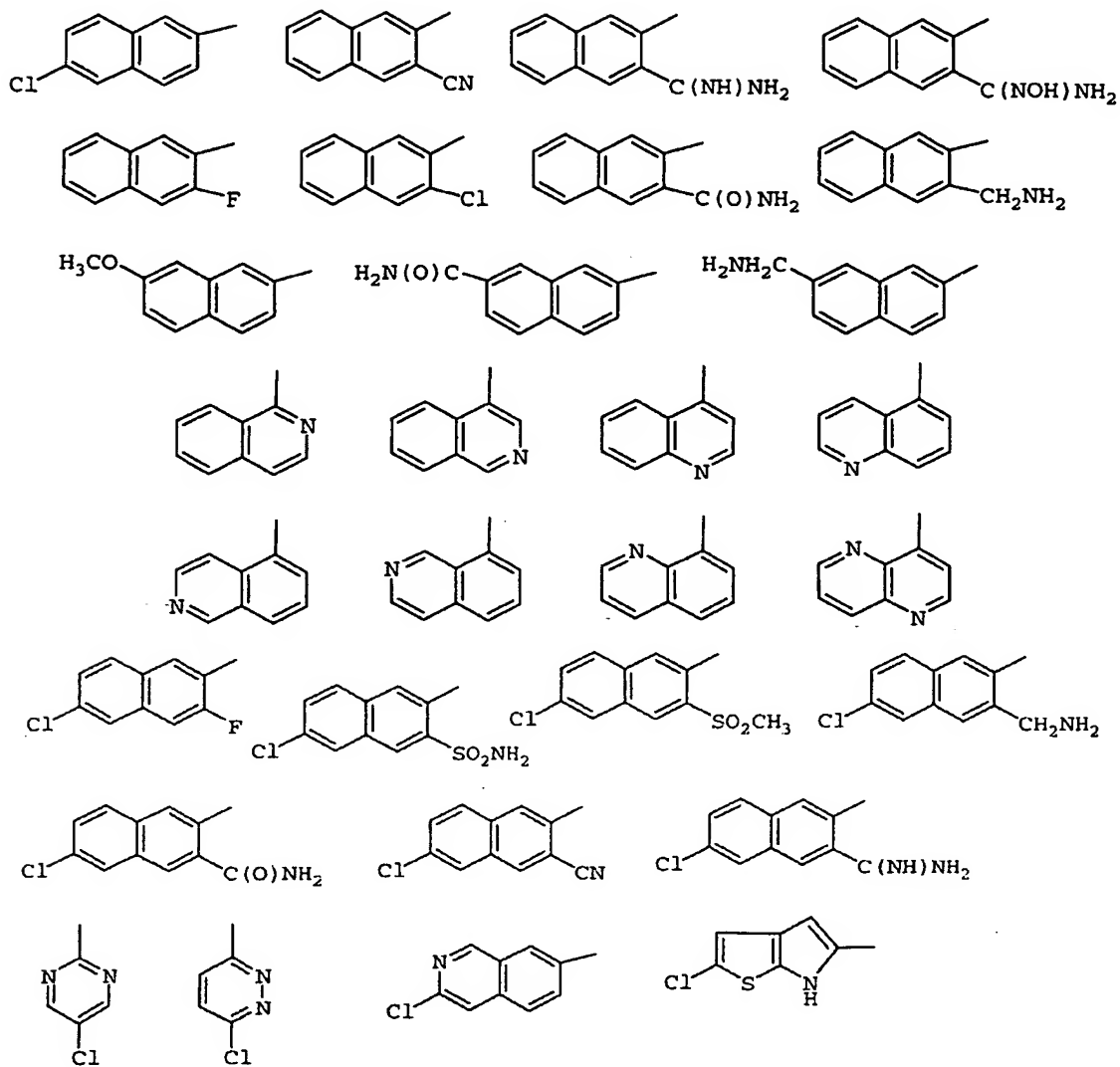
G is selected from the group:

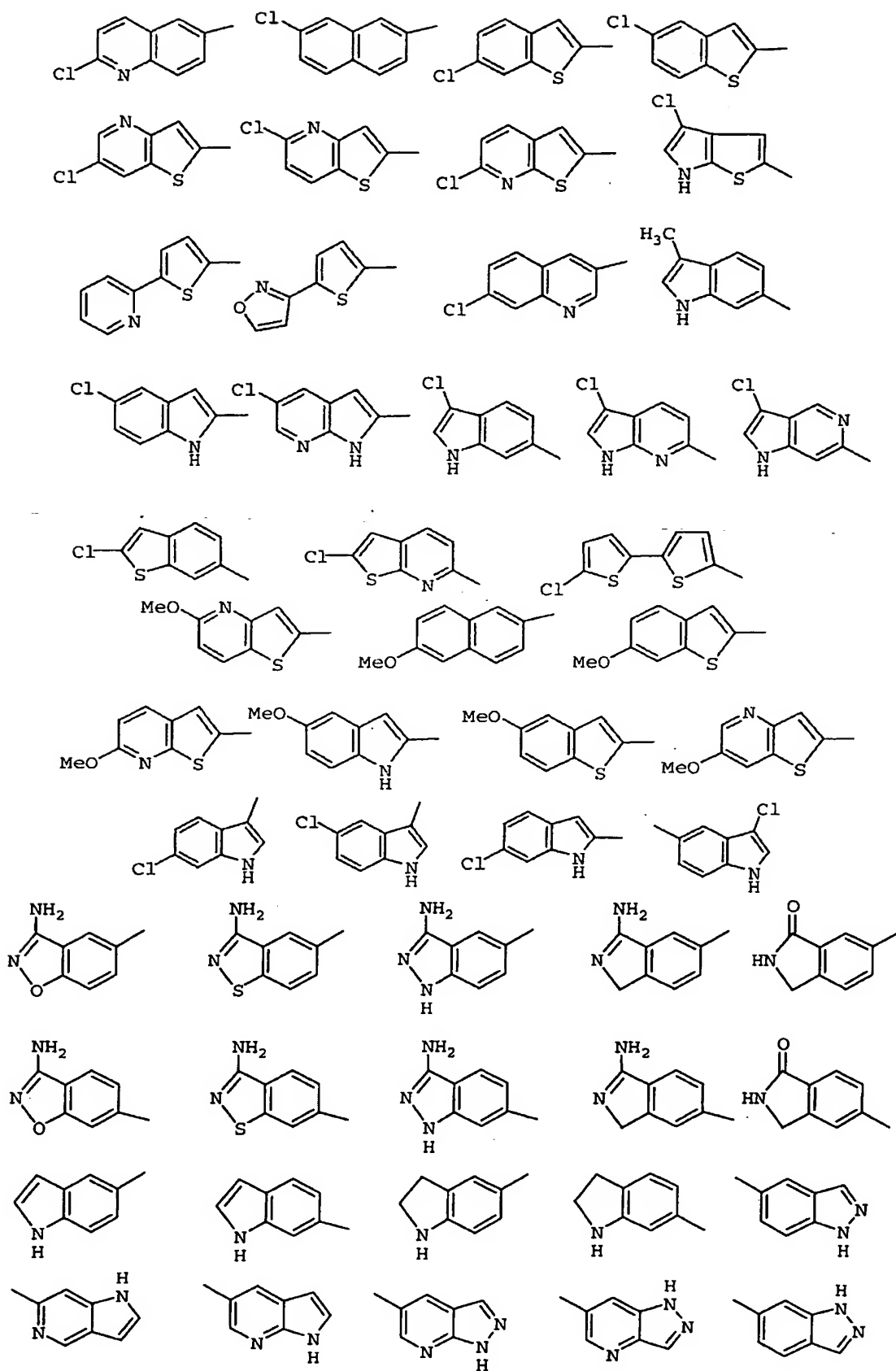
- phenyl; 2,5-bis-aminomethyl-phenyl;
- 5 2-amido-4-methoxy-phenyl; 2-amido-5-chloro-phenyl;
 - 2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
 - 2-aminomethyl-3-methoxy-phenyl;
 - 2-aminomethyl-4-fluoro-phenyl;
 - 2-aminomethyl-4-methoxy-phenyl;
 - 10 2-aminomethyl-5-fluoro-phenyl;
 - 2-aminomethyl-5-methoxy-phenyl;
 - 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 - 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
 - 2-aminosulfonyl-phenyl; 2-aminomethyl-4-ethyl-phenyl; 2-
 - 15 aminosulfonyl-4-ethyl-phenyl; 2-amido-4-ethyl-phenyl;
 - 2-hydroxy-4-methoxy-phenyl; 2-methylsulfonyl-phenyl;
 - 3-(N,N-dimethylamino)-4-chloro-phenyl;
 - 3-(N,N-dimethylamino)-phenyl; 3-(N-hydroxy-amidino)-phenyl;
 - 3-(N-methoxy-amidino)-phenyl;
 - 20 3-(N-methylamino)-4-chloro-phenyl;
 - 3-(N-methylamino)-phenyl; 3-amidino-phenyl;
 - 3-amido-6-hydroxy-phenyl; 3-amido-phenyl;
 - 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 - 3-amino-phenyl; 3-chloro-4-fluoro-phenyl; 3-chloro-phenyl;

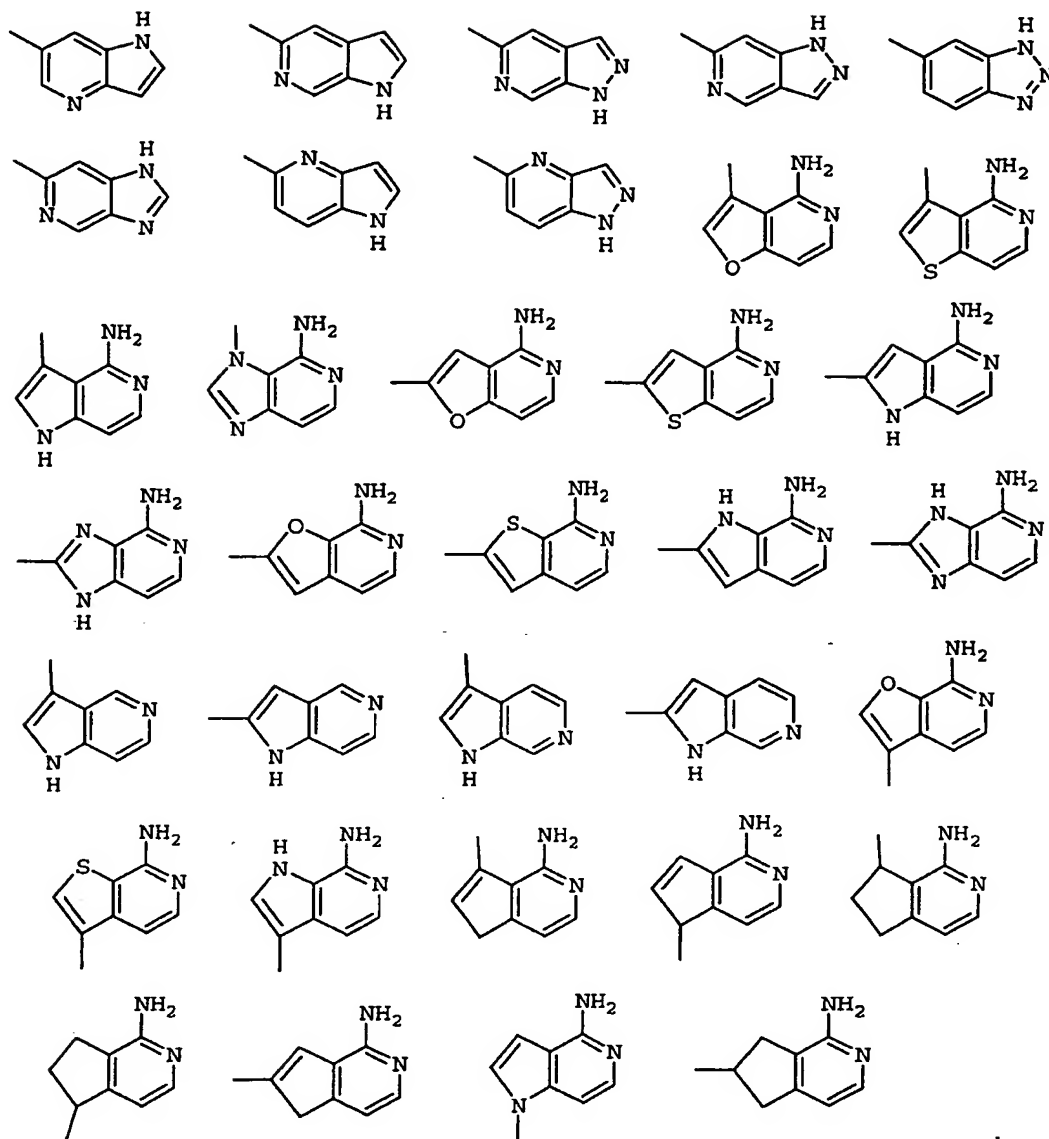
- 3-hydroxy-4-methoxy-phenyl;
 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
 4-(N-methylamino)-5-chloro-thien-2-yl;
 4-amino-5-chloro-thien-2-yl; 4-amino-pyrid-2-yl;
 5 4-chloro-3-fluoro-phenyl; 4-chloro-phenyl;
 4-chloro-pyrid-2-yl; 4-ethyl-phenyl; 4-ethyl-2-methylsulfonyl-phenyl; 4-ethyl-2-methoxy-phenyl;
 4-methoxy-2-methylsulfonyl-phenyl; 4-methoxy-phenyl;
 2-methoxy-pyrid-5-yl;
 10 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
 5-(N-methylamino)-4-chloro-thien-2-yl;
 5-amino-4-chloro-thien-2-yl;
 5-chloro-2-aminosulfonyl-phenyl;
 5-chloro-2-methylsulfonyl-phenyl; 5-chloro-pyrid-2-yl;
 15 5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
 5-methyl-thien-2-yl; 5-fluoro-thien-2-yl;
 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;











- 5 G_1 is absent or is selected from $(CR^3R^3a)_{1-3}$, $CR^3=CR^3$,
 $(CR^3R^3a)_u C(O)(CR^3R^3a)_w$, $(CR^3R^3a)_u O(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^3b(CR^3R^3a)_w$, $(CR^3R^3a)_u C(O)NR^3b(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^3bC(O)(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^3bC(O)(CR^3R^3a)_u C(O)NR^3b(CR^3R^3a)_w$,
10 $(CR^3R^3a)_u S(CR^3R^3a)_w$, $(CR^3R^3a)_u S(O)(CR^3R^3a)_w$,
 $(CR^3R^3a)_u S(O)_2(CR^3R^3a)_w$, $(CR^3R^3a)_u S(O)NR^3b(CR^3R^3a)_w$,
 $(CR^3R^3a)_u NR^3bS(O)_2(CR^3R^3a)_w$, and
 $(CR^3R^3a)_u S(O)_2NR^3b(CR^3R^3a)_w$, wherein $u + w$ total 0, 1,

or 2, provided that G₁ does not form a N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

- 5 A is selected from one of the following carbocycles and heterocycles which are substituted with 0-2 R⁴;
 cyclohexyl, phenyl, piperidiny1, piperazinyl, pyridyl, pyrimidyl, furanyl, morpholinyl, thienyl, pyrrolyl, pyrrolidinyl, oxazolyl, isoxazolyl,
 10 thiazolyl, isothiazolyl, pyrazolyl, imidazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl,
 15 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl, 1,3,4-triazolyl, benzofuranyl, benzothiofuranyl, indolinyl, indolyl, benzimidazolyl, benzoxazolyl, benzthiazolyl, indazolyl, benzisoxazolyl, benzisothiazolyl, and isoindazolyl;

20

X is selected from -(CR²R^{2a})₁₋₂-, -C(O)-, -S(O)₂-, -NR²S(O)₂-, -NR²S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR², -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

- 25 Y is a C₃₋₆ monocyclic carbocycle or 5-6 membered monocyclic heterocycle, wherein the carbocycle or heterocycle consists of carbon atoms and 0-2 heteroatoms selected from N, O, and S(O)_p, the carbocycle or heterocycle further comprises 0-1 double bonds and 0-1 carbonyl groups, and the carbocycle or heterocycle is
 30 substituted with 0-2 R⁴;

alternatively, Y is CY¹Y², and Y¹ and Y² are independently C₁₋₂ alkyl substituted with 0-1 R⁴;

35

- R^{1a} , at each occurrence, is selected from H, R^{1b} , $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, CH_2R^{1b} , and $CH_2CH_2R^{1b}$, provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;
- 5 alternatively, when two R^{1a} groups are attached to adjacent atoms or to the same carbon atom, together with the atoms to which they are attached they form a 5-6 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N, 10 O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b} and 0-3 ring double bonds;

- R^{1b} is selected from H, CH_3 , CH_2CH_3 , F, Cl, Br, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , 15 $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, phenyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 20 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

- R^2 , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 25 0-2 R^{4b} , benzyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

- 30 R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, phenyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon

atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom
5 to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

10

R^{2b}, at each occurrence, is selected from CF₃, C₁₋₄ alkoxy, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-2 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2c}, at each occurrence, is selected from CF₃, OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃,
20 CH(CH₃)₂, benzyl, phenyl substituted with 0-2 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

25

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4
30 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group

consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

- 5 R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4
 10 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms
 15 other than a C(O)-halo or C(O)-S(O)_p moiety;

R⁴, at each occurrence, is selected from H, (CH₂)₂OR², CH₂OR², OR², F, Cl, Br, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃,
 20 C(CH₃)₃, -CN, NO₂, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, C(O)R^{2c}, NR²C(O)R^{2b}, C(O)NR²R^{2a}, SO₂NR²R^{2a}, CF₃, and CF₂CF₃;

R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle
 25 substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d}, (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d},
 30 (CR³R^{3g})_r-NR^{2d}C(O)R^{2e}, (CR³R^{3g})_r-C(O)R^{2e}, (CR³R^{3g})_r-OC(O)R^{2e}, (CR³R^{3g})_r-C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-C(O)OR^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)OR^{2d}, (CR³R^{3g})_r-SO₂NR^{2d}R^{2d},

$(\text{CR}^3\text{R}^3\text{g})_r\text{-NR}^{2d}\text{SO}_2\text{R}^{2d}$, and $(\text{CR}^3\text{R}^3\text{g})_r\text{-S(O)}_p\text{R}^{2d}$, provided that $\text{S(O)}_p\text{R}^{2d}$ forms other than $\text{S(O)}_2\text{H}$ or S(O)H ;

R^{4b} , at each occurrence, is selected from H, =O, OR^3 ,
 5 CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, -CN,
 NO_2 , NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, C(O)R^3 , $\text{CH}_2\text{-C(O)R}^3$, C(O)OR^{3c} ,
 $\text{CH}_2\text{-C(O)OR}^{3c}$, $\text{NR}^3\text{C(O)R}^{3a}$, $\text{CH}_2\text{NR}^3\text{C(O)R}^{3a}$, $\text{C(O)NR}^3\text{R}^{3a}$, $\text{CH}_2\text{-}$
 $\text{C(O)NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{CH}_2\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl,
 $\text{CH}_2\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl, $\text{NR}^3\text{SO}_2\text{-phenyl}$, $\text{CH}_2\text{NR}^3\text{SO}_2\text{-phenyl}$,
 10 $\text{S(O)}_p\text{CF}_3$, $\text{CH}_2\text{S(O)}_p\text{CF}_3$, $\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $\text{CH}_2\text{S(O)}_p\text{-C}_{1-4}$
 alkyl, $\text{S(O)}_p\text{-phenyl}$, $\text{CH}_2\text{S(O)}_p\text{-phenyl}$, and CF_3 ;

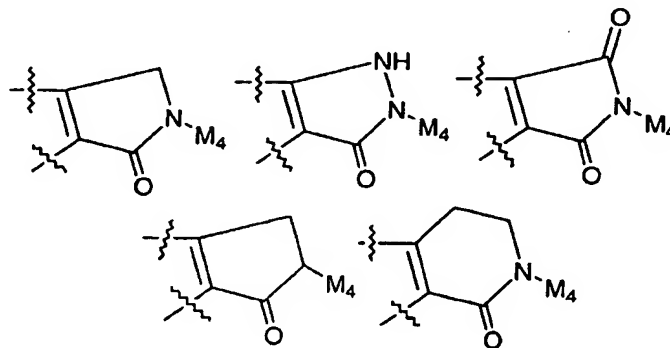
R^{4c} , at each occurrence, is selected from =O, OR^2 ,
 $(\text{CR}^3\text{R}^{3a})\text{OR}^2$, F, $(\text{CR}^3\text{R}^{3a})\text{F}$, Br, $(\text{CR}^3\text{R}^{3a})\text{Br}$, Cl,
 15 $(\text{CR}^3\text{R}^{3a})\text{Cl}$, CF_3 , $(\text{CR}^3\text{R}^{3a})\text{CF}_3$, C_{1-4} alkyl, C_{2-3} alkenyl,
 C_{2-3} alkynyl, -CN, $(\text{CR}^3\text{R}^{3a})\text{CN}$, NO_2 , $(\text{CR}^3\text{R}^{3a})\text{NO}_2$, $\text{NR}^{2a}\text{R}^{2a}$,
 $(\text{CR}^3\text{R}^{3a})\text{NR}^{2a}\text{R}^{2a}$, $\text{N(}\rightarrow\text{O)R}^{2a}\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{N(}\rightarrow\text{O)R}^{2a}\text{R}^{2a}$, C(O)R^{2c} ,
 $(\text{CR}^3\text{R}^{3a})\text{C(O)R}^{2c}$, $\text{NR}^{2c}\text{C(O)R}^{2b}$, $(\text{CR}^3\text{R}^{3a})\text{NR}^{2c}\text{C(O)R}^{2b}$,
 $\text{C(O)NR}^{2a}\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{C(O)NR}^{2a}\text{R}^{2a}$, $\text{NR}^{2c}\text{C(O)NR}^{2a}\text{R}^{2a}$,
 20 $(\text{CR}^3\text{R}^{3a})\text{NR}^{2c}\text{C(O)NR}^{2a}\text{R}^{2a}$, $\text{SO}_2\text{NR}^{2a}\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{SO}_2\text{NR}^{2a}\text{R}^{2a}$,
 $\text{NR}^{2a}\text{SO}_2\text{NR}^{2a}\text{R}^{2a}$, $(\text{CR}^3\text{R}^{3a})\text{NR}^{2a}\text{SO}_2\text{NR}^{2a}\text{R}^{2a}$, $\text{NR}^{2a}\text{SO}_2\text{R}^{5a}$,
 $(\text{CR}^3\text{R}^{3a})\text{NR}^{2a}\text{SO}_2\text{R}^{5a}$, $\text{S(O)}_p\text{R}^{5a}$, $(\text{CR}^3\text{R}^{3a})\text{S(O)}_p\text{R}^{5a}$, CF_3 ,
 CF_2CF_3 , C_{3-10} carbocycle substituted with 0-2 R^{4b} ,
 $(\text{CR}^3\text{R}^{3a})\text{-C}_{3-10}$ carbocycle substituted with 0-2 R^{4b} , 5-10
 25 membered heterocycle substituted with 0-2 R^{4b} and
 consisting of carbon atoms and from 1-4 heteroatoms
 selected from the group consisting of N, O, and S(O)_p ,
 and $(\text{CR}^3\text{R}^{3a})\text{-5-10}$ membered heterocycle substituted with
 0-2 R^{4b} and consisting of carbon atoms and from 1-4
 30 heteroatoms selected from the group consisting of N,
 O, and S(O)_p ;

- R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$,
 $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -phenyl, $S(O)_pCF_3$,
 $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 , phenyl substituted
 with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
 benzyl substituted with 0-2 R^6 ; and,
 R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN, NO_2 , NR^2R^{2a} ,
 $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$, $SO_2NR^2R^{2a}$,
 and $NR^2SO_2C_{1-4}$ alkyl.

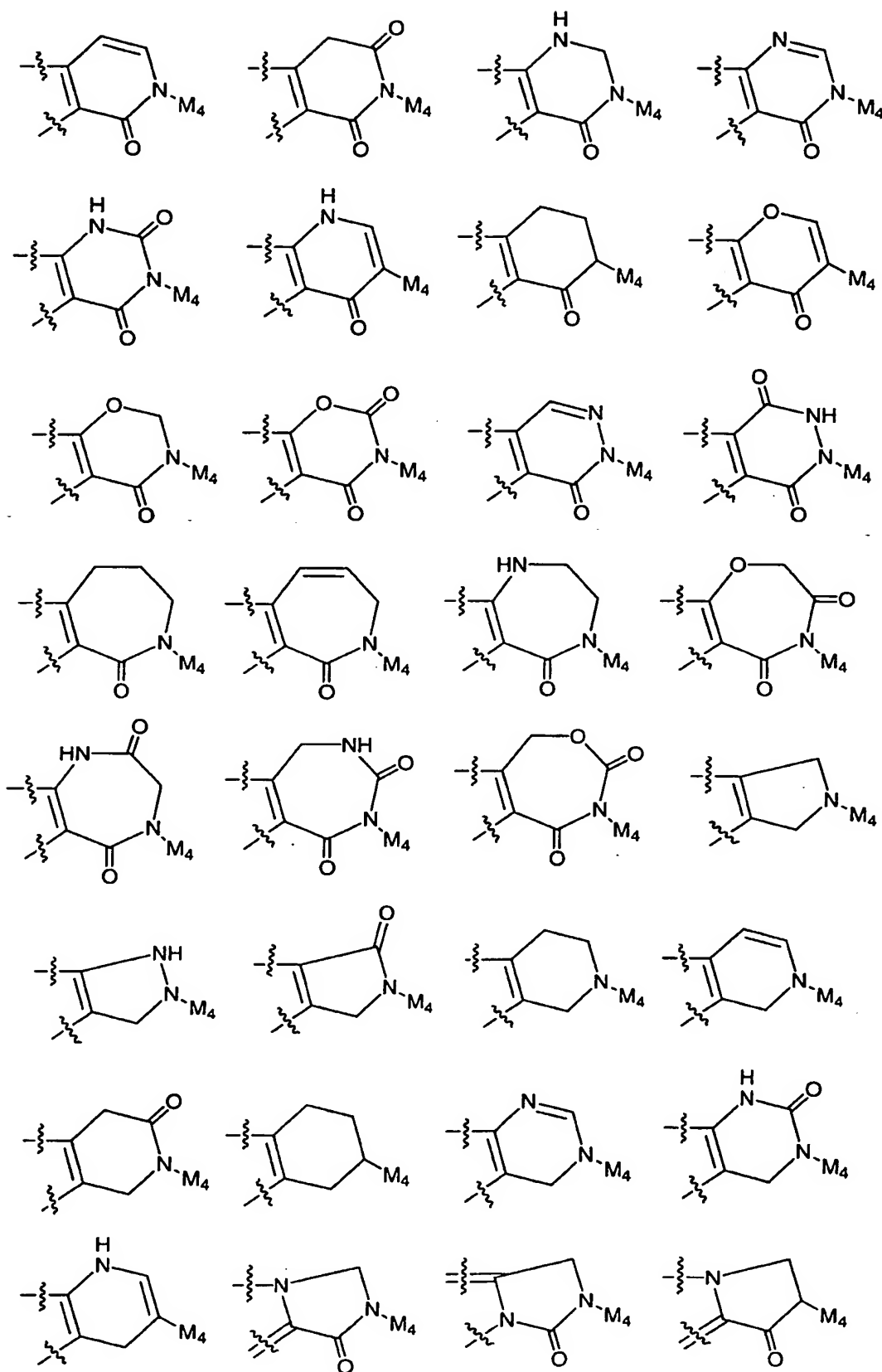
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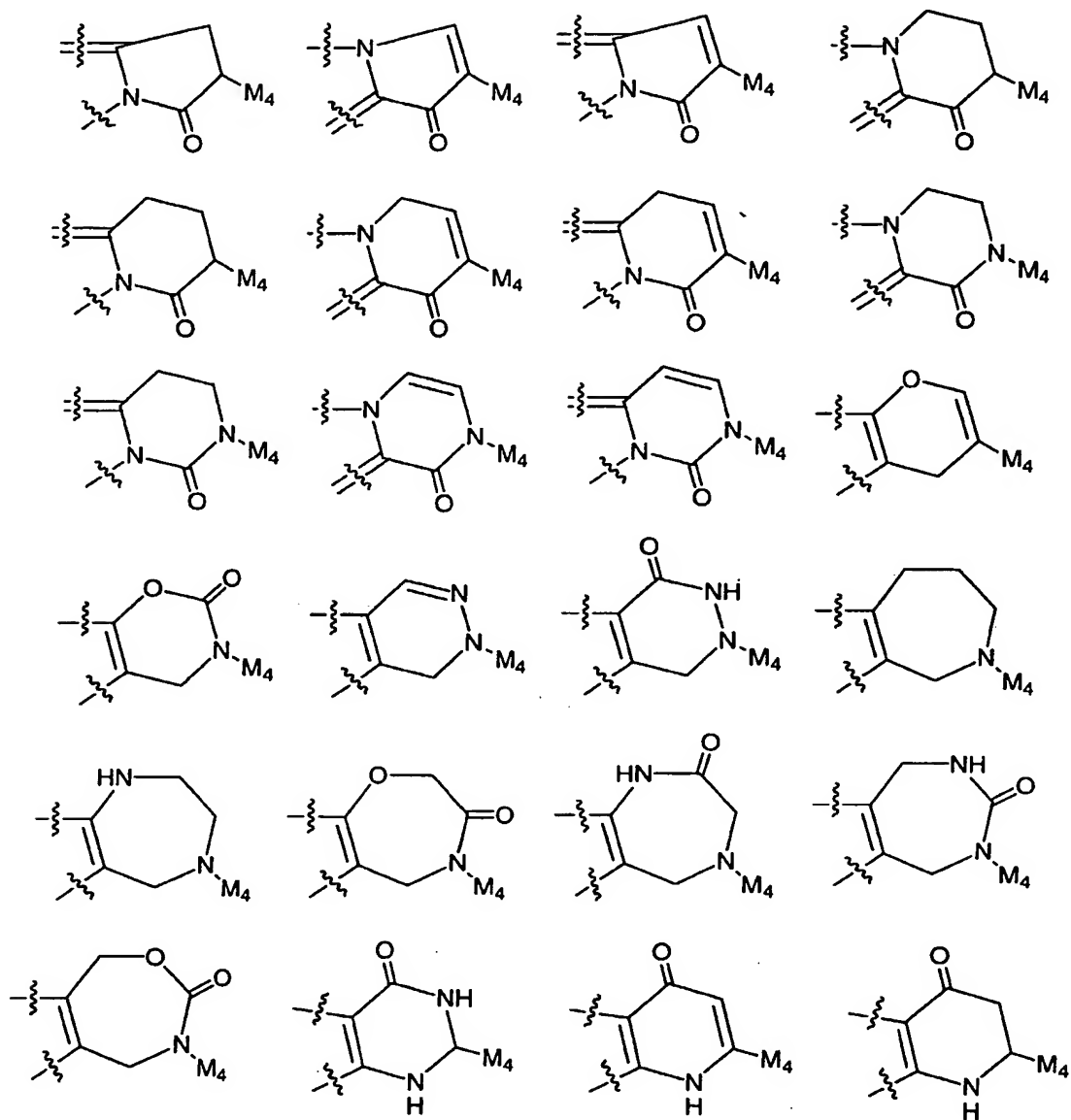
4. A compound according to Claim 3, wherein:

ring M is substituted with 0-2 R^{1a} and is selected from the group:

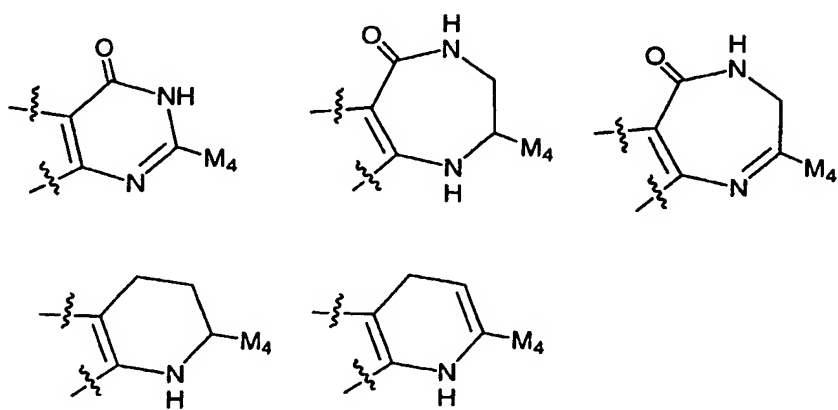


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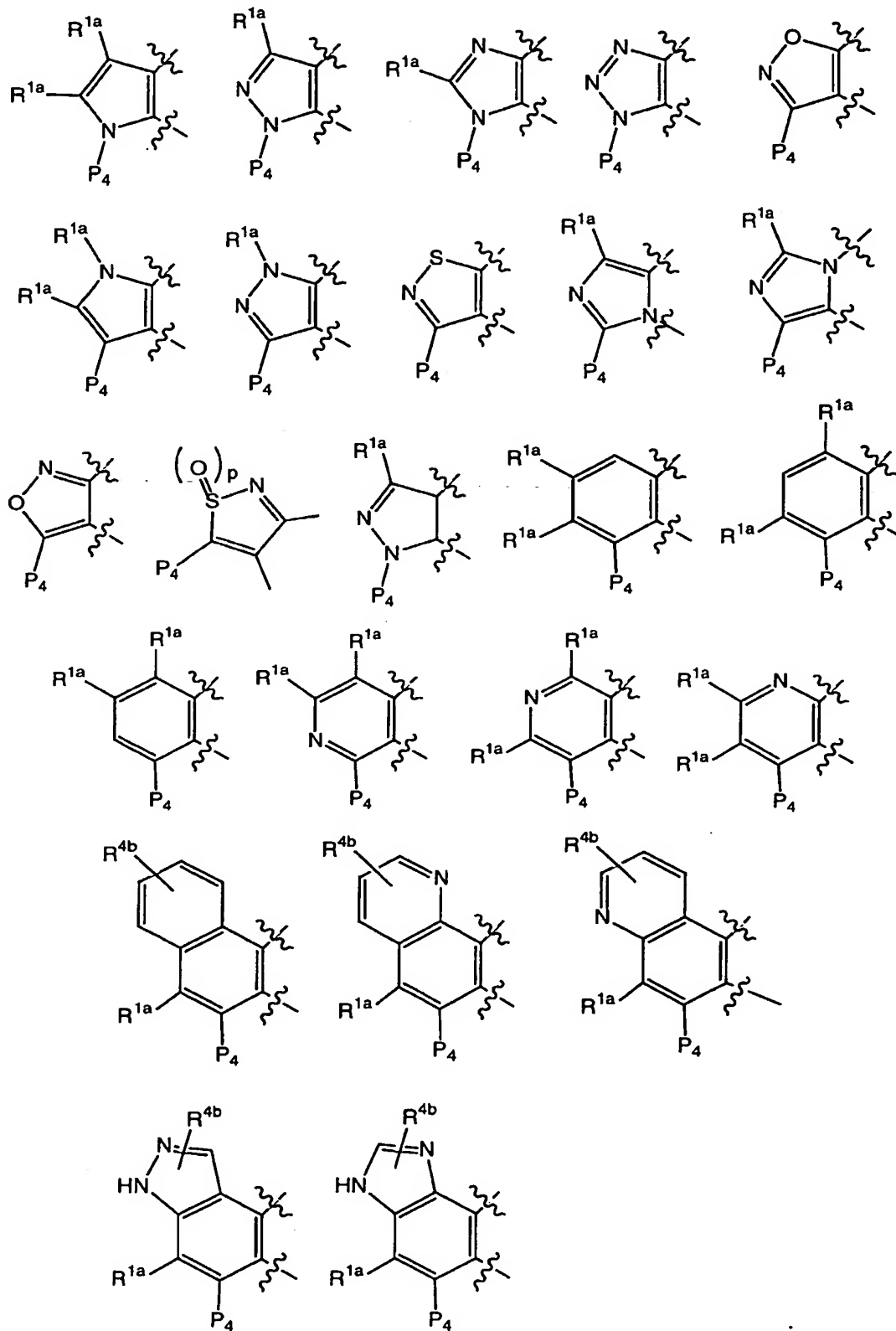


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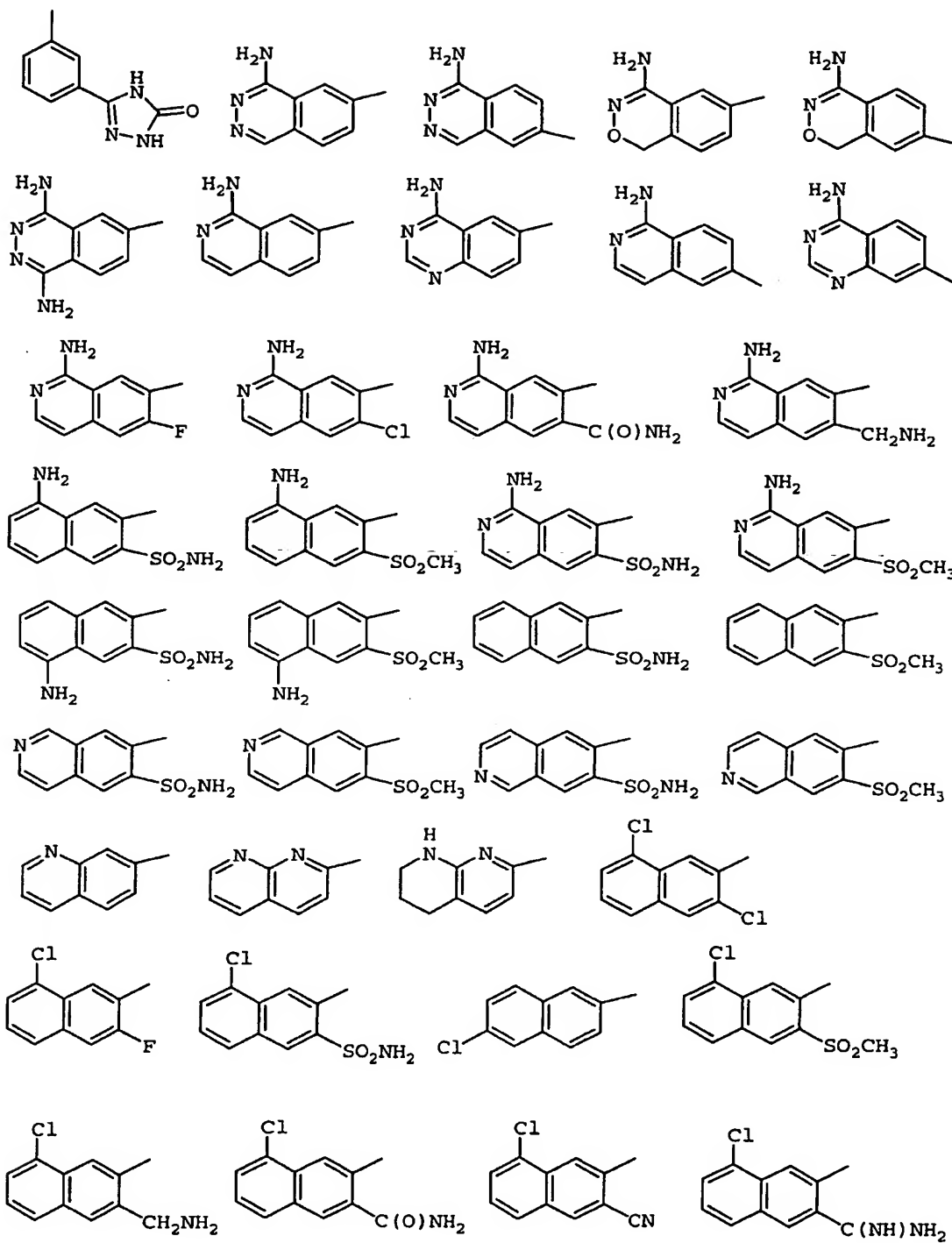
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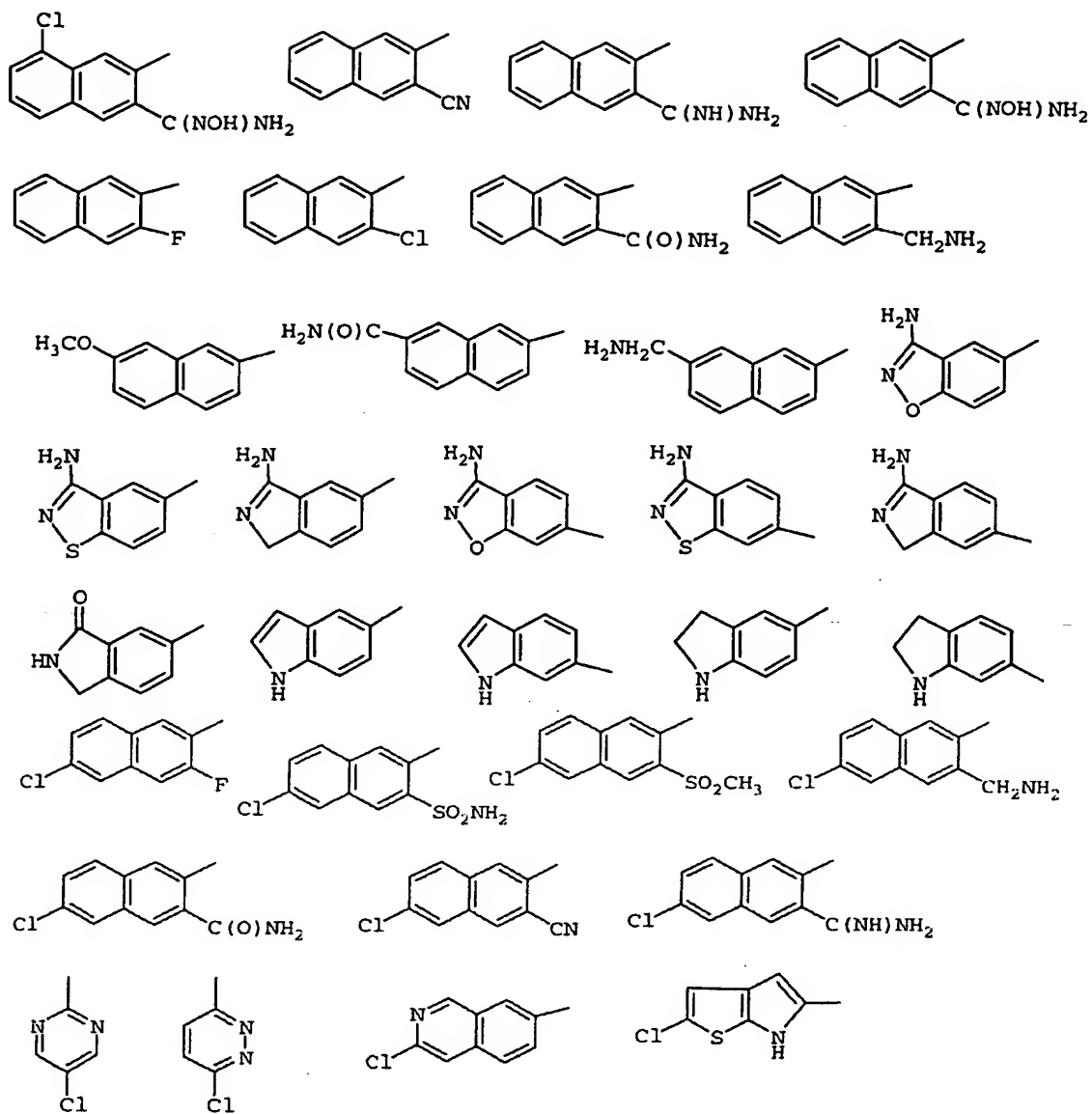
ring P, including P₁, P₂, P₃, and P₄ is selected from group:

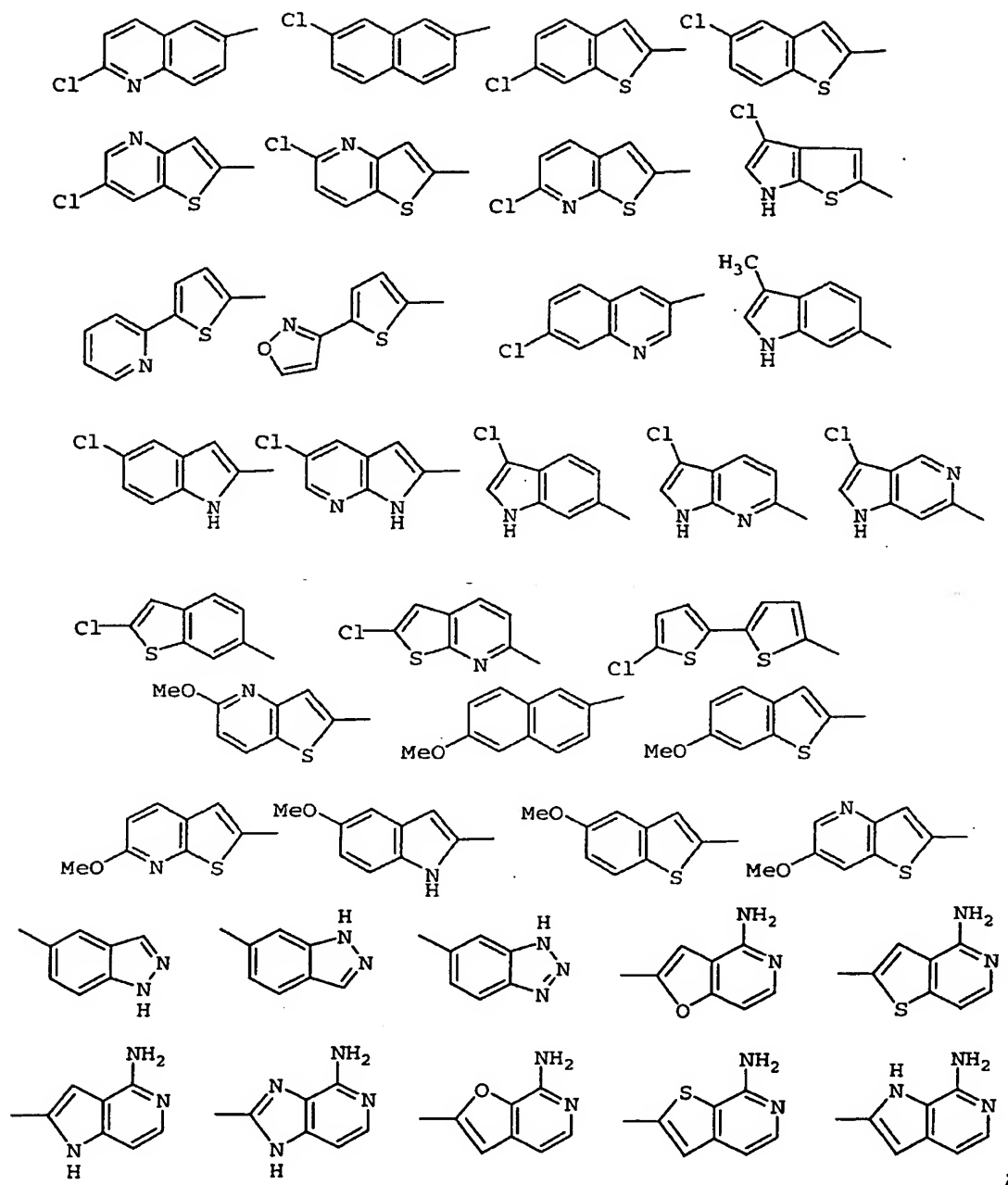


G is selected from the group:

- phenyl; 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
2-aminomethyl-3-fluoro-phenyl;
5 2-aminomethyl-4-fluoro-phenyl;
2-aminomethyl-4-methoxy-phenyl;
2-aminomethyl-5-fluoro-phenyl;
2-aminomethyl-5-methoxy-phenyl;
2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
10 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
2-aminosulfonyl-phenyl; 2-methylsulfonyl-phenyl; 2-
aminomethyl-4-ethyl-phenyl; 2-aminosulfonyl-4-ethyl-phenyl;
2-amido-4-ethyl-phenyl;
3-(N,N-dimethylamino)-4-chloro-phenyl;
15 3-(N,N-dimethylamino)-phenyl;
3-(N-methylamino)-4-chloro-phenyl;
3-(N-methylamino)-phenyl; 3-amido-phenyl;
3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
3-amino-phenyl; 3-chloro-phenyl;
20 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
4-(N-methylamino)-5-chloro-thien-2-yl;
4-amino-5-chloro-thien-2-yl; 4-chloro-phenyl; 4-ethyl-
phenyl; 4-ethyl-2-methylsulfonyl-phenyl; 4-ethyl-2-methoxy-
phenyl; 4-methoxy-2-methylsulfonyl-phenyl;
25 4-methoxy-phenyl;
5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
5-(N-methylamino)-4-chloro-thien-2-yl;
5-amino-4-chloro-thien-2-yl; 5-chloro-pyrid-2-yl;
5-chloro-thien-2-yl; 5-methoxy-thien-2-yl;
30 5-methyl-thien-2-yl; 5-fluoro-thien-2-yl;
6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;







- 5 G_1 is absent or is selected from CH_2 , CH_2CH_2 , $CH=CH$, CH_2O , OCH_2 , NH , CH_2NH , $NHCH_2$, $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $CH_2S(O)_2$, $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$, provided that G_1 does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;

- A is selected from cyclohexyl, piperidinyl, phenyl, pyridyl, and pyrimidyl, and is substituted with 0-2 R^4 ;
- 5 X is selected from CH_2 , $C(O)$, $-S(O)_2-$, $-NHC(O)-$, $-C(O)NH-$, $-CH_2NH-$, O, and $-CH_2O-$;
- Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentanonyl, cyclohexyl, cyclohexanonyl, pyrrolidinyl, pyrrolidinonyl, 10 piperidinyl, piperidinonyl, tetrahydrofuranyl, and tetrahydropyranyl, and, when Y is a ring, Y is substituted with 0-1 R^4 ;
- 15 R^{1a} , at each occurrence, is selected from H, R^{1b} , $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, and CH_2R^{1b} , provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;
- R^{1b} is selected from CH_3 , CH_2CH_3 , F, Cl, Br, $-CN$, CF_3 , OR^2 , 20 NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , CO_2R^{2a} , $S(O)_pR^2$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided 25 that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;
- R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-1 R^{4b} , 30 benzyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2a}, at each occurrence, is selected from H, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2b}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2c}, at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, benzyl, phenyl substituted with 0-1 R^{4b}, and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4

heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted with 0-2 R^{4c}, -(CH₂)-C₃₋₆ carbocycle substituted with 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, and -(CH₂)-5-6 membered heterocycle and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

R⁴, at each occurrence, is selected from OH, OR², CH₂OR², (CH₂)₂OR², F, Br, Cl, I, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, CF₃, and CF₂CF₃;

R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d}, (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d}, (CR³R^{3g})_r-C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)R^{2e},

$(CR^3R^3g)_x-C(O)R^{2e}$, $(CR^3R^3g)_x-NR^{2d}C(O)NR^{2d}R^{2d}$,
 $(CR^3R^3g)_x-NR^{2d}C(O)OR^{2d}$, $(CR^3R^3g)_x-NR^{2d}SO_2R^{2d}$, and
 $(CR^3R^3g)_x-S(O)_pR^{2d}$, provided that $S(O)_pR^{2d}$ forms other
than $S(O)_2H$ or $S(O)H$;

5

R^{4b} , at each occurrence, is selected from H, =O, OR^3 ,
 CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN,
 NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $C(O)OR^{3c}$, $NR^3C(O)R^{3a}$,
 $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, NR^3SO_2 -phenyl,
10 $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, and CF_3 ;

R^{4c} , at each occurrence, is selected from =O, OR^2 , CH_2OR^2 ,
F, Br, Cl, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, C_{2-3}
alkenyl, C_{2-3} alkynyl, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$,
15 $N(\rightarrow O)R^2R^{2a}$, $CH_2N(\rightarrow O)R^2R^{2a}$, $C(O)R^{2c}$, $CH_2C(O)R^{2c}$,
 $NR^2C(O)R^{2b}$, $CH_2NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $CH_2C(O)NR^2R^{2a}$,
 $SO_2NR^2R^{2a}$, $CH_2SO_2NR^2R^{2a}$, $NR^2SO_2R^{5a}$, $CH_2NR^2SO_2R^{5a}$,
 $S(O)_pR^{5a}$, $CH_2S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , C_{3-6} carbocycle
substituted with 0-2 R^{4b} , $(CH_2)-C_{3-6}$ carbocycle
20 substituted with 0-2 R^{4b} , 5-6 membered heterocycle
substituted with 0-2 R^{4b} and consisting of carbon atoms
and from 1-4 heteroatoms selected from the group
consisting of N, O, and $S(O)_p$, and $(CH_2)-5-6$ membered
heterocycle substituted with 0-2 R^{4b} and consisting of
25 carbon atoms and from 1-4 heteroatoms selected from
the group consisting of N, O, and $S(O)_p$;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
30 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $C(O)OR^{3c}$, $NR^3C(O)R^{3a}$,
 $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, NR^3SO_2 -phenyl,
 $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 , phenyl substituted

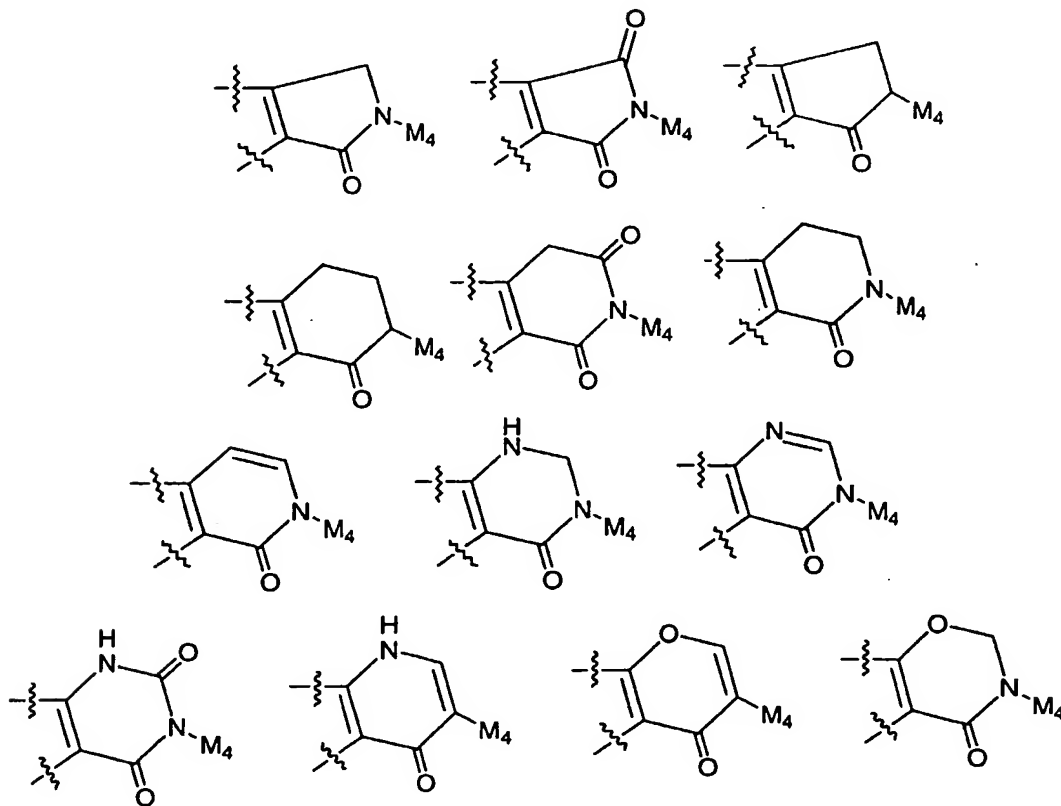
with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
benzyl substituted with 0-2 R^6 ;

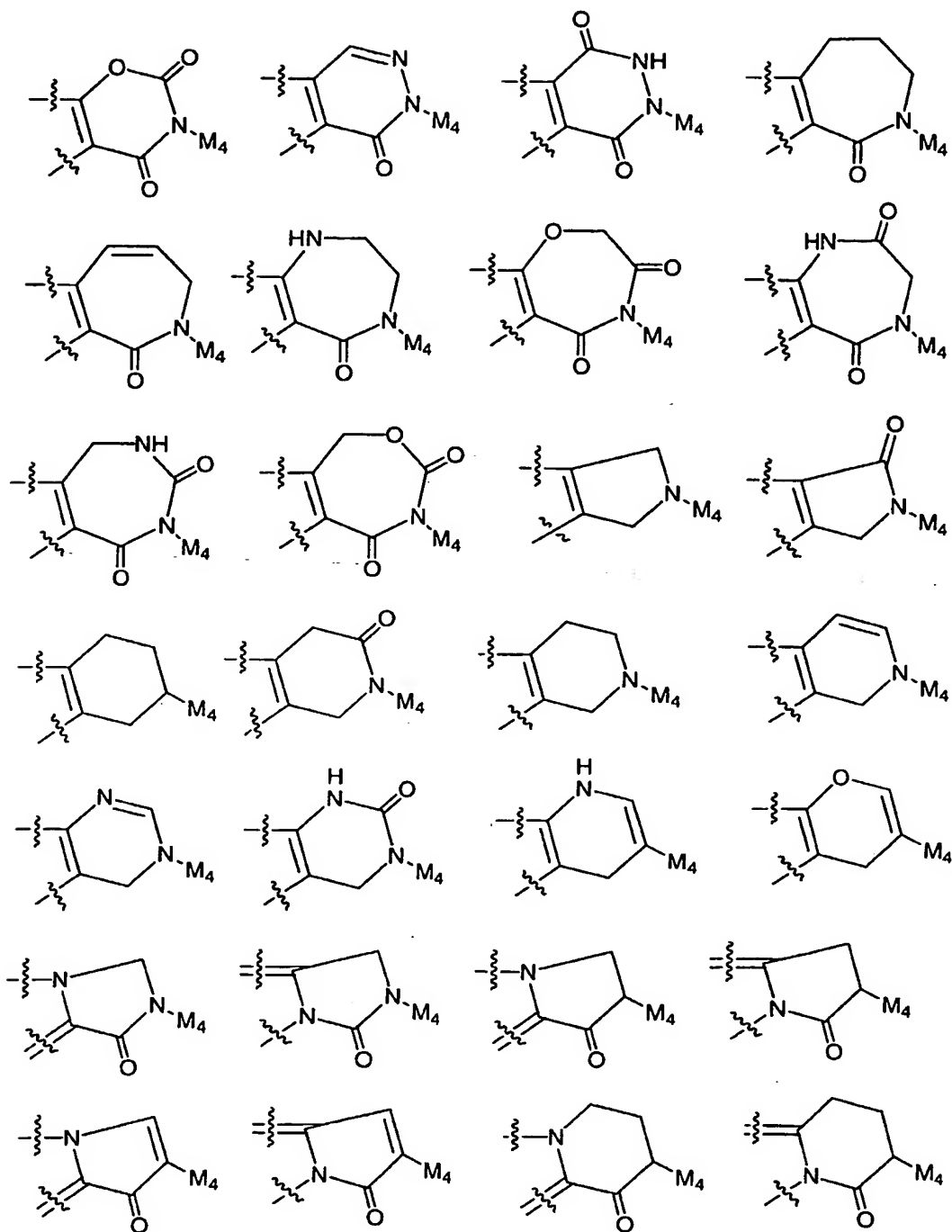
R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
5 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $-CN$, NO_2 , NR^2R^{2a} ,
 $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$, and
 $SO_2NR^2R^{2a}$; and,

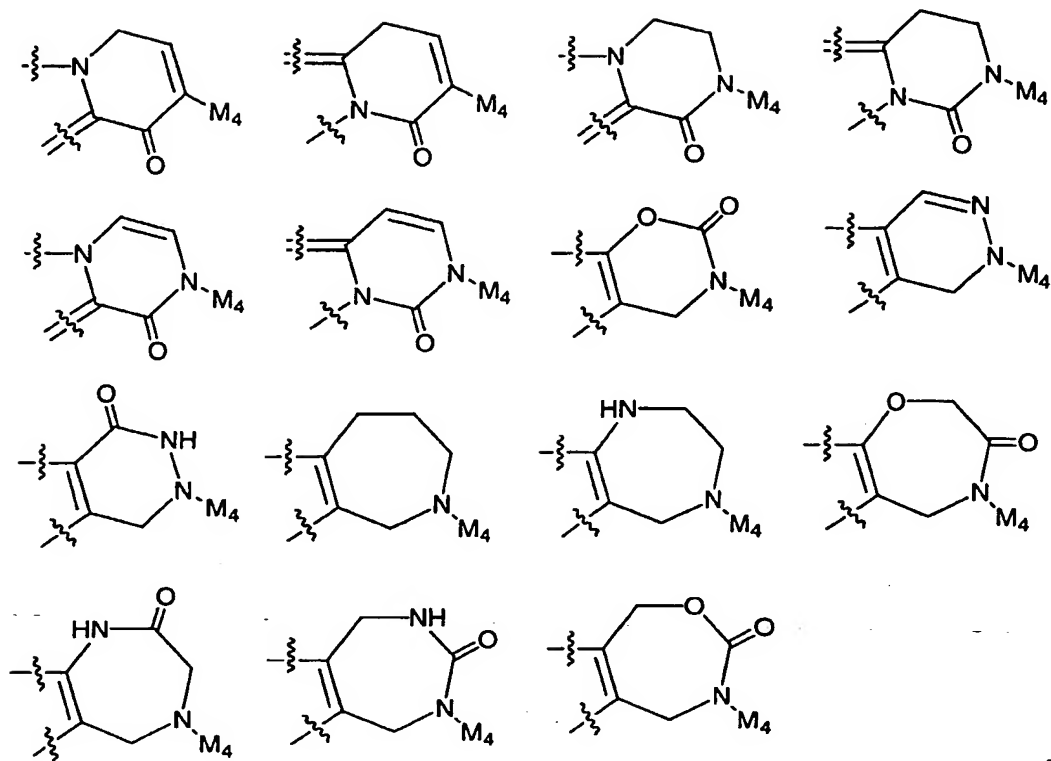
r , at each occurrence, is selected from 0, 1, and 2.
10

5. A compound according to Claim 4, wherein:

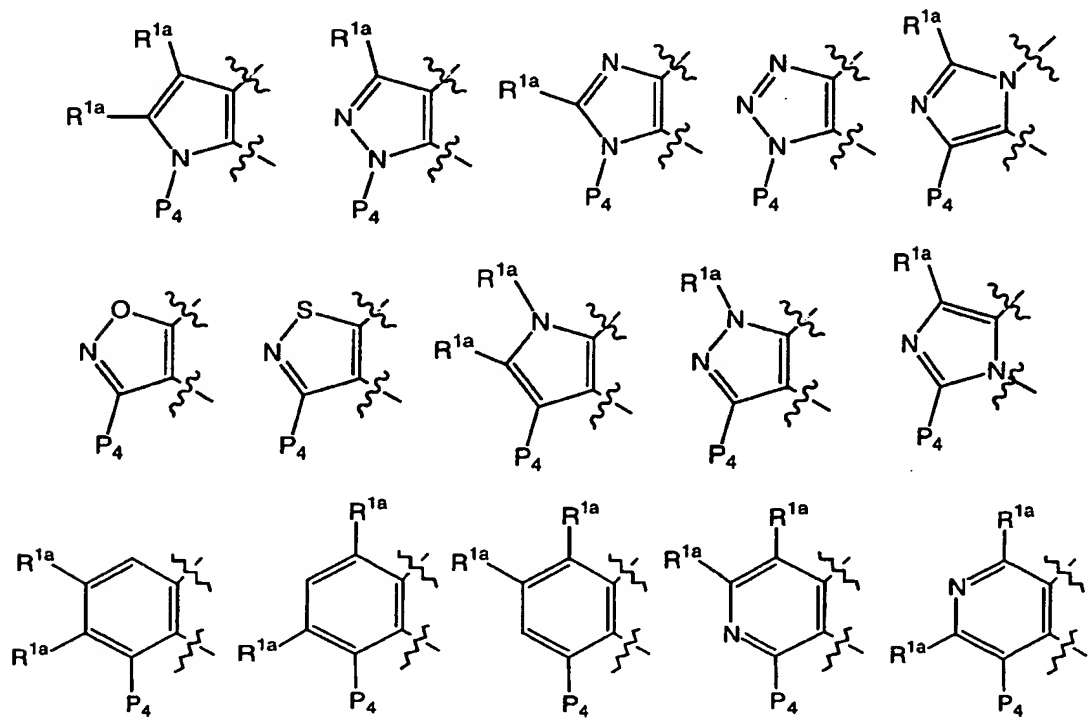
15 ring M is substituted with 0-1 R^{1a} and is selected from the
group:







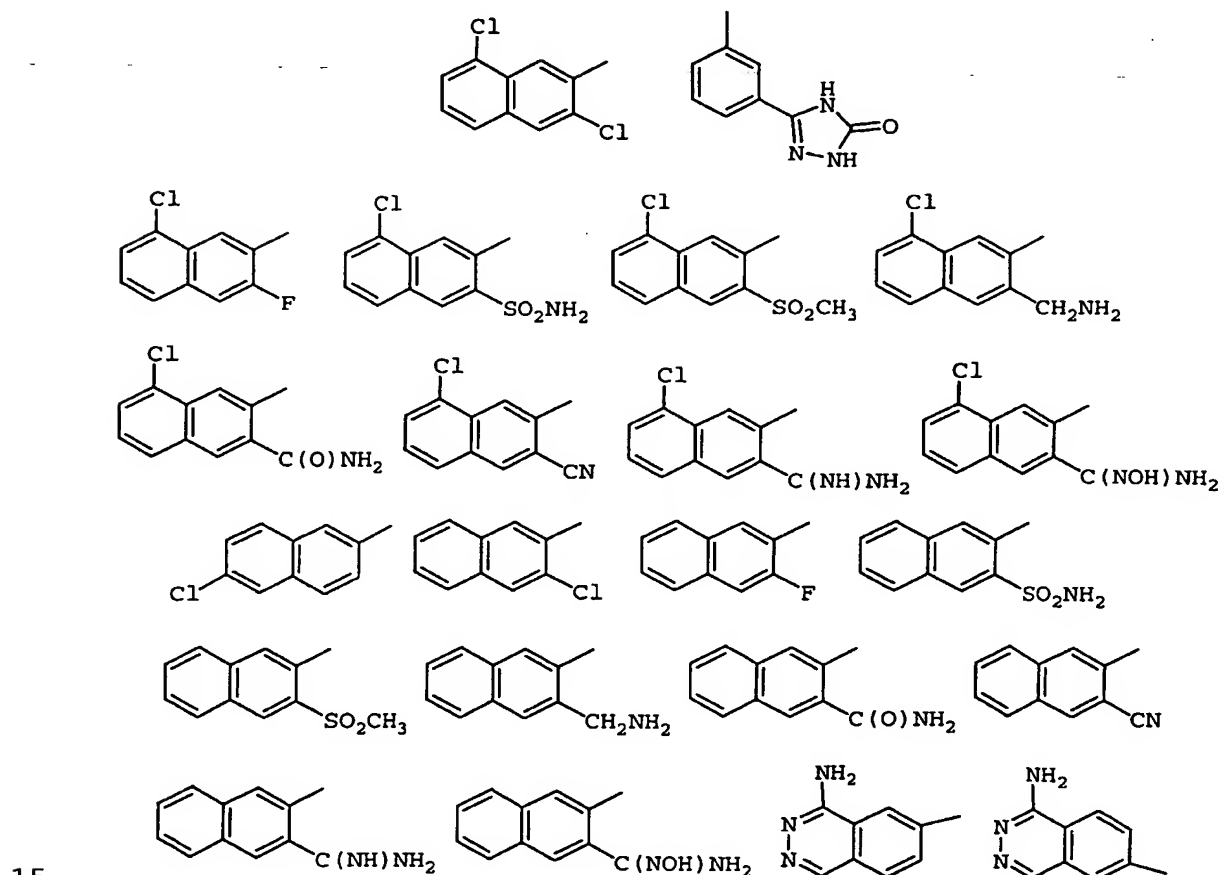
ring P, including P_1 , P_2 , P_3 , and P_4 is selected from group:

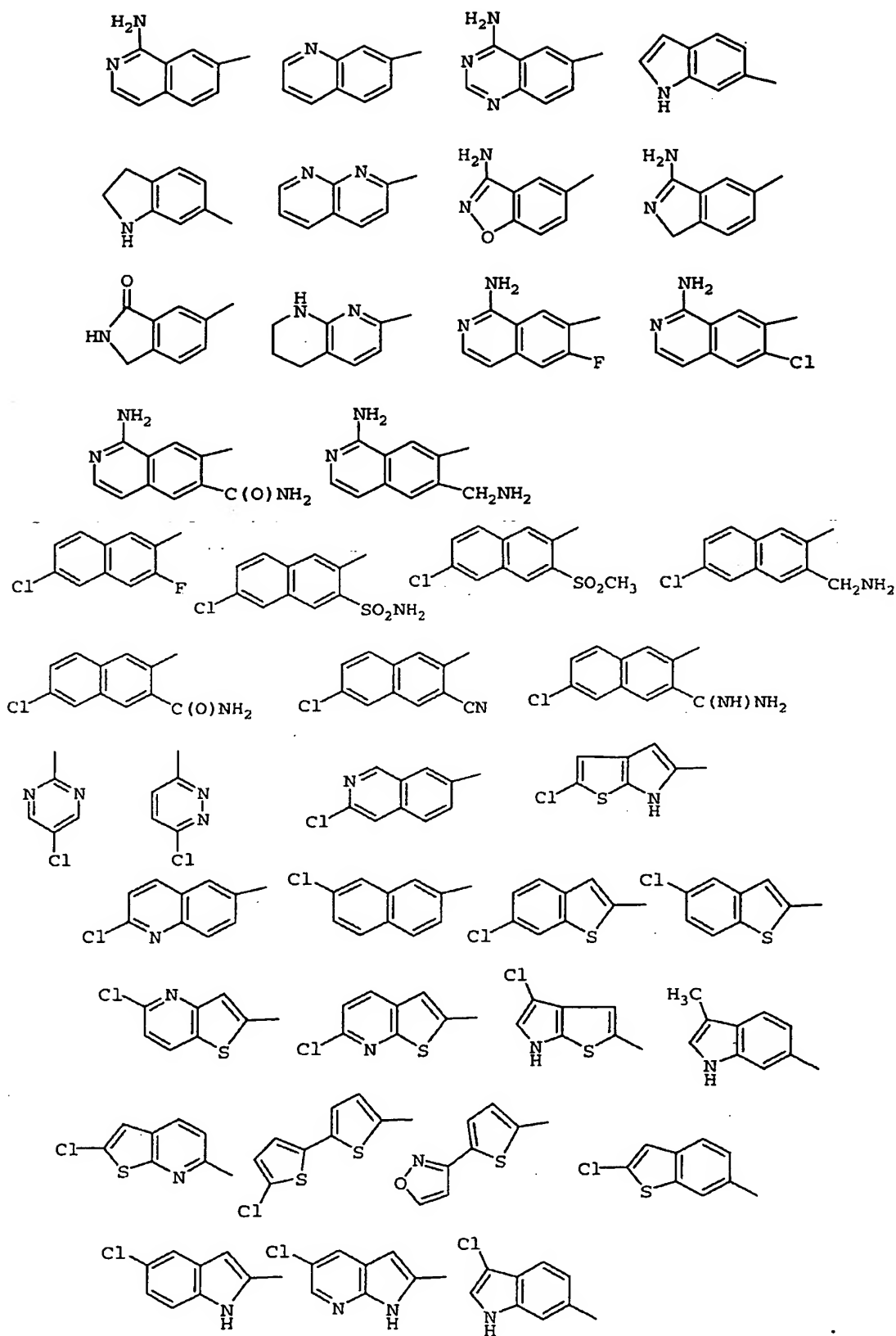


5

-G is selected from:

- 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
 2-aminomethyl-3-fluoro-phenyl;
 2-aminomethyl-4-fluoro-phenyl;
 5 2-aminomethyl-5-fluoro-phenyl;
 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
 2-aminosulfonyl-phenyl; 3-amido-phenyl;
 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 10 3-chloro-phenyl; 4-chloro-phenyl; 4-ethyl-phenyl;
 4-methoxy-phenyl; 5-chloro-pyrid-2-yl; 5-chloro-thien-2-yl;
 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl;





A is selected from the group: cyclohexyl, piperidinyl, phenyl, 2-pyridyl, 3-pyridyl, 2-pyrimidyl, 2-Cl-phenyl, 3-Cl-phenyl, 2-F-phenyl, 3-F-phenyl, 2-methylphenyl, 2-aminophenyl, and 2-methoxyphenyl;

5

Y is selected from C(CH₃)₂, C(CH₂CH₃)₂, cyclopropyl, cyclobutyl, cyclopentyl, 2-cyclopentanonyl, cyclohexyl, 2-cyclohexanonyl, pyrrolidinyl (attached to A and R^{4a} at the 2-position), pyrrolidinyl (attached to A and R^{4a} at the 3-position), 2-pyrrolidinonyl (attached to A and R^{4a} at the 3-position), piperidinyl (attached to A and R^{4a} at the 4-position), 4-piperidinonyl (attached to A and R^{4a} at the 3-position), tetrahydrofuranyl, and tetrahydropyranyl (attached to A and R^{4a} at the 4-position);

10

15

R^{1a}, at each occurrence, is selected from H, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH₂F, CH₂Cl, Br, CH₂Br, -CN, CH₂CN, CF₃, CH₂CF₃, OCH₃, CH₂OH, C(CH₃)₂OH, CH₂OCH₃, NH₂, CH₂NH₂, NHCH₃, CH₂NHCH₃, N(CH₃)₂, CH₂N(CH₃)₂, CO₂H, COCH₃, CO₂CH₃, CH₂CO₂CH₃, SCH₃, CH₂SCH₃, S(O)CH₃, CH₂S(O)CH₃, S(O)₂CH₃, CH₂S(O)₂CH₃, C(O)NH₂, CH₂C(O)NH₂, SO₂NH₂, CH₂SO₂NH₂, NHSO₂CH₃, CH₂NHSO₂CH₃, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyridin-2-yl-N-oxide, pyridin-3-yl-N-oxide, pyridin-4-yl-N-oxide, imidazol-1-yl, CH₂-imidazol-1-yl, 4-methyl-oxazol-2-yl, 4-N,N-dimethylaminomethyl-oxazol-2-yl, 1,2,3,4-tetrazol-1-yl, 1,2,3,4-tetrazol-5-yl, CH₂-1,2,3,4-tetrazol-1-yl, and CH₂-1,2,3,4-tetrazol-5-yl, provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

20

25

30

R², at each occurrence, is selected from H, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, phenyl substituted with 0-1 R^{4b}, benzyl substituted with 0-1 R^{4b}, and 5 membered

aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

- 5 R^{2a} , at each occurrence, is selected from H, CH₃, and CH₂CH₃;

alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6
10 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

- 15 R^{2b} , at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, CH₃, and CH₂CH₃;

R^{2c} , at each occurrence, is selected from OH, OCH₃, OCH₂CH₃, CH₃, and CH₂CH₃;

20

- R^{2d} , at each occurrence, is selected from H, R^{4c} , C₁₋₄ alkyl substituted with 0-2 R^{4c} , C₃₋₆ cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6
25 membered aromatic heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

30

R^{2e} , at each occurrence, is selected from H, R^{4c} , C₁₋₄ alkyl substituted with 0-2 R^{4c} , C₃₋₆ cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c}

and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

5

R^{4a} is selected from -(CH₂)_r-5-6 membered carbocycle substituted with 0-3 R^{4c}, -(CH₂)_r-5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, (CH₂)_rNR^{2d}R^{2d}, (CH₂)_rN(→O)R^{2d}R^{2d}, (CH₂)_rOR^{2d}, (CH₂)_r-C(O)NR^{2d}R^{2d}, (CH₂)_r-NR^{2d}C(O)R^{2e}, (CH₂)_r-C(O)R^{2e}, (CH₂)_r-NR^{2d}C(O)NR^{2d}R^{2d}, (CH₂)_r-NR^{2d}C(O)OR^{2d}, (CH₂)_r-NR^{2d}SO₂R^{2d}, and (CH₂)_r-S(O)_pR^{2d}, provided that S(O)_pR^{2d} forms other than S(O)₂H or S(O)H;

10

15

R^{4b}, at each occurrence, is selected from H, =O, OR³, CH₂OR³, F, Cl, CH₃, CH₂CH₃, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, C(O)OR^{3c}, NR³C(O)R^{3a}, C(O)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂-phenyl, S(O)₂CH₃, S(O)₂-phenyl, and CF₃;

20

R^{4c}, at each occurrence, is selected from =O, OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, C₂₋₃ alkenyl, C₂₋₃ alkynyl, CH₂OH, CH₂OCH₃, CH₂OCH₂CH₃, CH₂OCH₂CH₂CH₃, CH₂OCH(CH₃)₂, F, Br, Cl, CF₃, NR²R^{2a}, CH₂NR²R^{2a}, N(→O)R²R^{2a}, CH₂N(→O)R²R^{2a}, C(O)R^{2c}, CH₂C(O)R^{2c}, NR²C(O)R^{2b}, CH₂NR²C(O)R^{2b}, C(O)NR²R^{2a}, CH₂C(O)NR²R^{2a}, SO₂NR²R^{2a}, CH₂SO₂NR²R^{2a}, NR²SO₂R^{5a}, CH₂NR²SO₂R^{5a}, S(O)_pR^{5a}, CH₂S(O)_pR^{5a}, CF₃, cyclopropyl substituted with 0-1 R^{4b}, cyclobutyl substituted with 0-1 R^{4b}, cyclopentyl substituted with 0-1 R^{4b}, phenyl substituted with 0-1 R^{4b}, -CH₂-cyclopropyl substituted

25

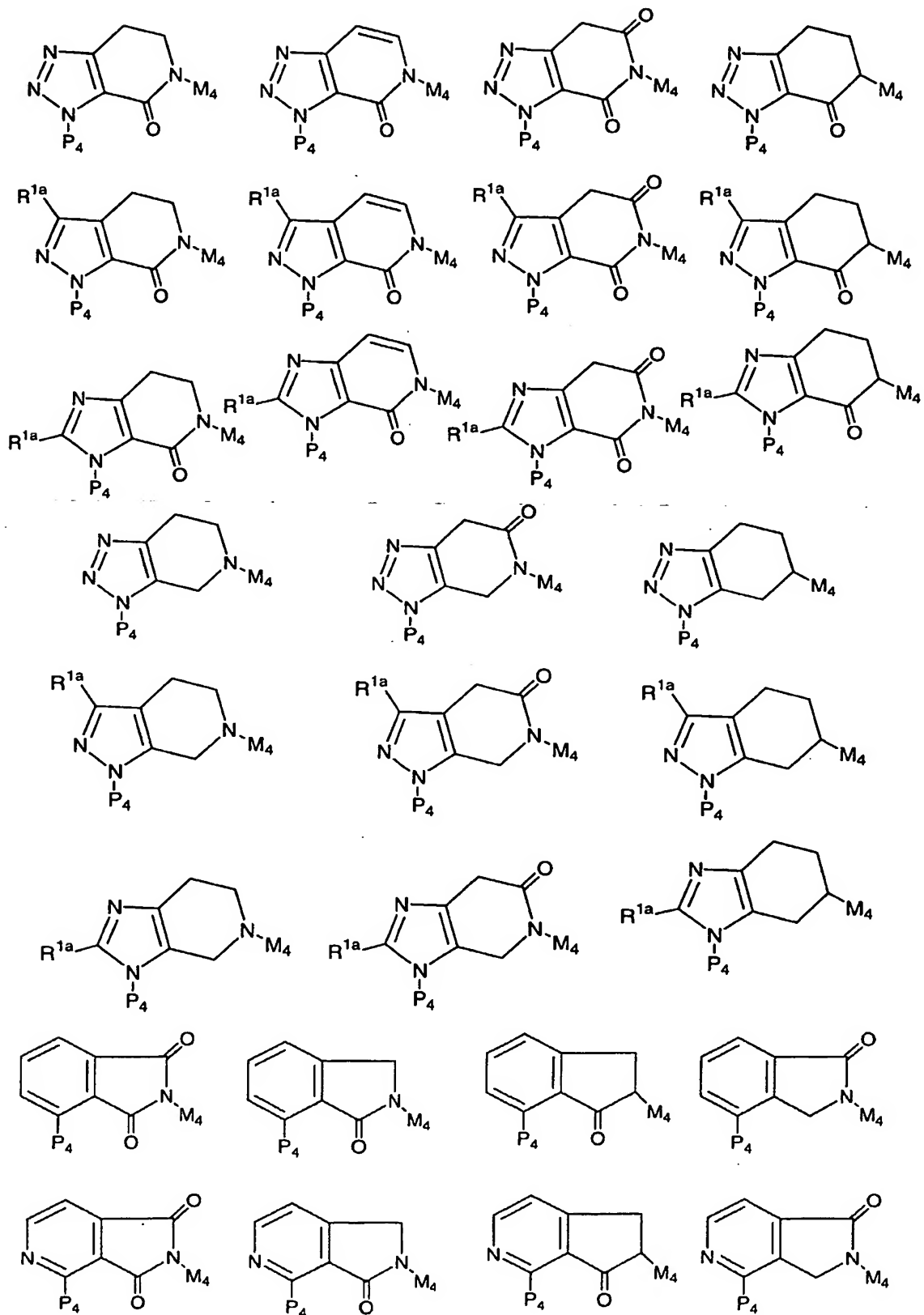
30

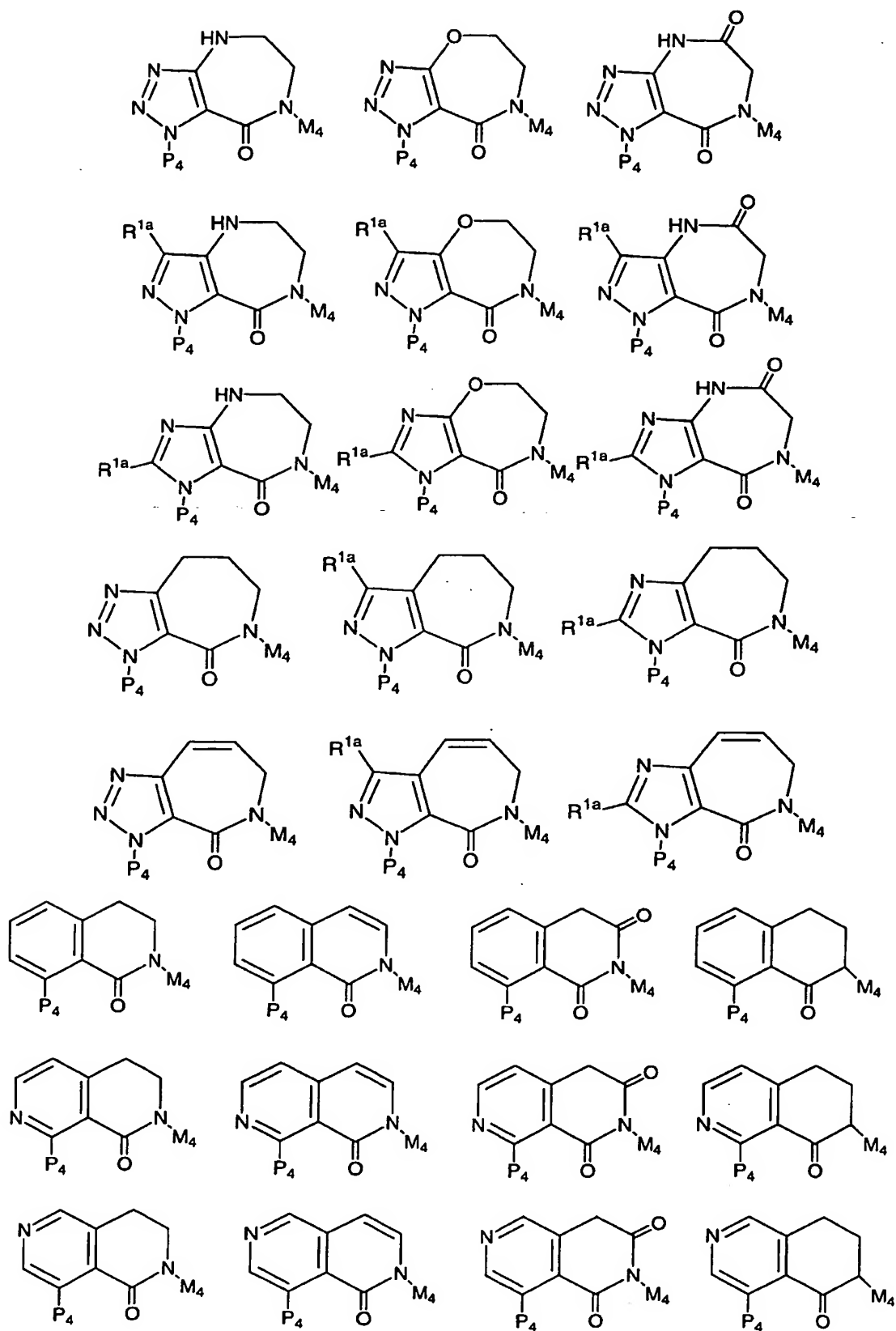
with 0-1 R^{4b} , $-\text{CH}_2$ -cyclobutyl substituted with 0-1 R^{4b} ,
- CH_2 -cyclopentyl substituted with 0-1 R^{4b} , benzyl
substituted with 0-2 R^{4b} , 5-6 membered aromatic
heterocycle substituted with 0-2 R^{4b} and consisting of
5 carbon atoms and from 1-4 heteroatoms selected from
the group consisting of N, O, and $\text{S}(\text{O})_p$, and $(\text{CH}_2)_{5-6}$
membered aromatic heterocycle substituted with 0-2 R^{4b}
and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
10 O, and $\text{S}(\text{O})_p$;

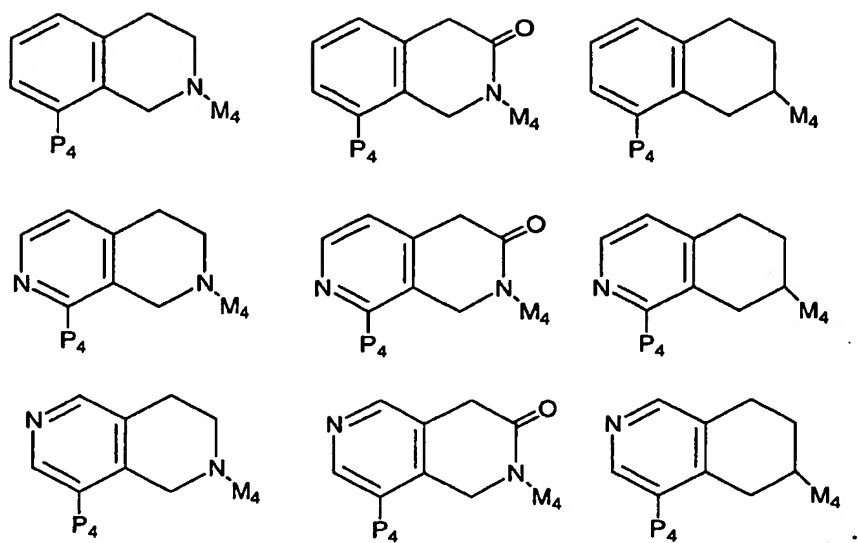
R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 OR^3 , CH_2OR^3 , F, Cl, NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, $\text{C}(\text{O})\text{R}^3$, $\text{C}(\text{O})\text{OR}^{3c}$,
 $\text{NR}^3\text{C}(\text{O})\text{R}^{3a}$, $\text{C}(\text{O})\text{NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl,
15 NR^3SO_2 -phenyl, $\text{S}(\text{O})_2\text{-CH}_3$, $\text{S}(\text{O})_2$ -phenyl, CF_3 , phenyl
substituted with 0-2 R^6 , naphthyl substituted with 0-2
 R^6 , and benzyl substituted with 0-2 R^6 ; and,

R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
20 CH_3 , CH_2CH_3 , NR^2R^{2a} , $\text{CH}_2\text{NR}^2\text{R}^{2a}$, $\text{C}(\text{O})\text{R}^{2b}$, $\text{CH}_2\text{C}(\text{O})\text{R}^{2b}$,
 $\text{NR}^2\text{C}(\text{O})\text{R}^{2b}$, and $\text{SO}_2\text{NR}^2\text{R}^{2a}$.

25 6. A compound according to Claim 5, wherein the
compound is selected from:





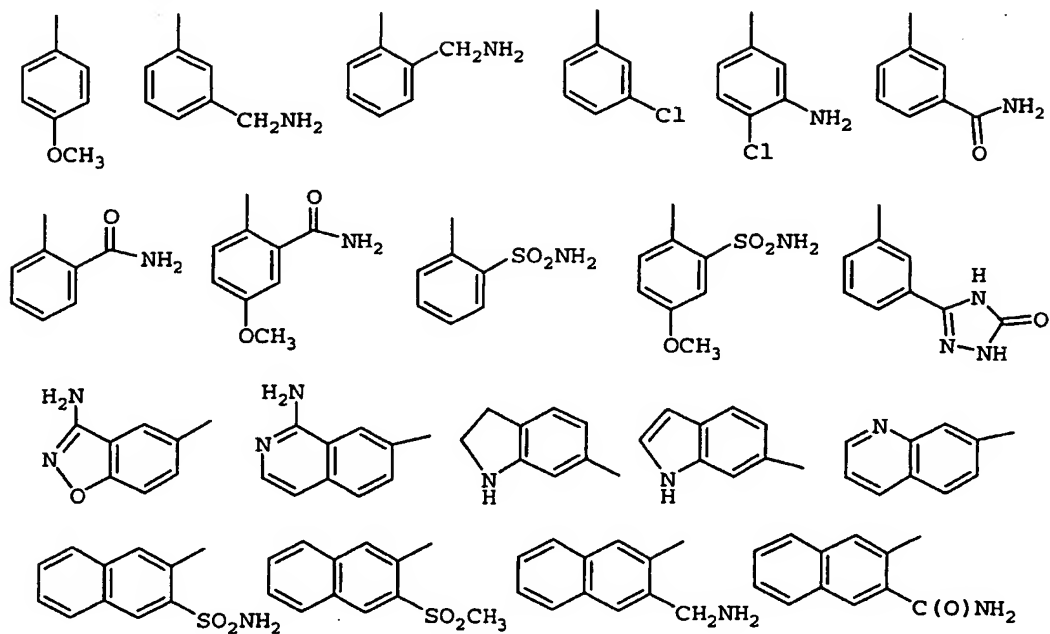


P_4 is $-G_1-G$;

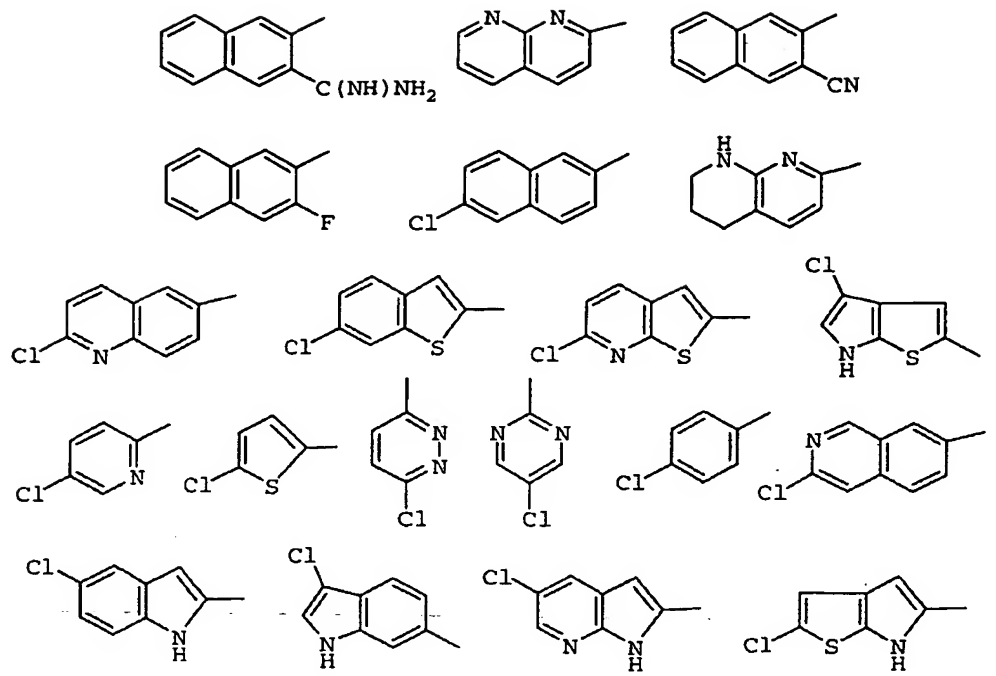
5

M_4 is $-A-B$;

$-G$ is selected from:



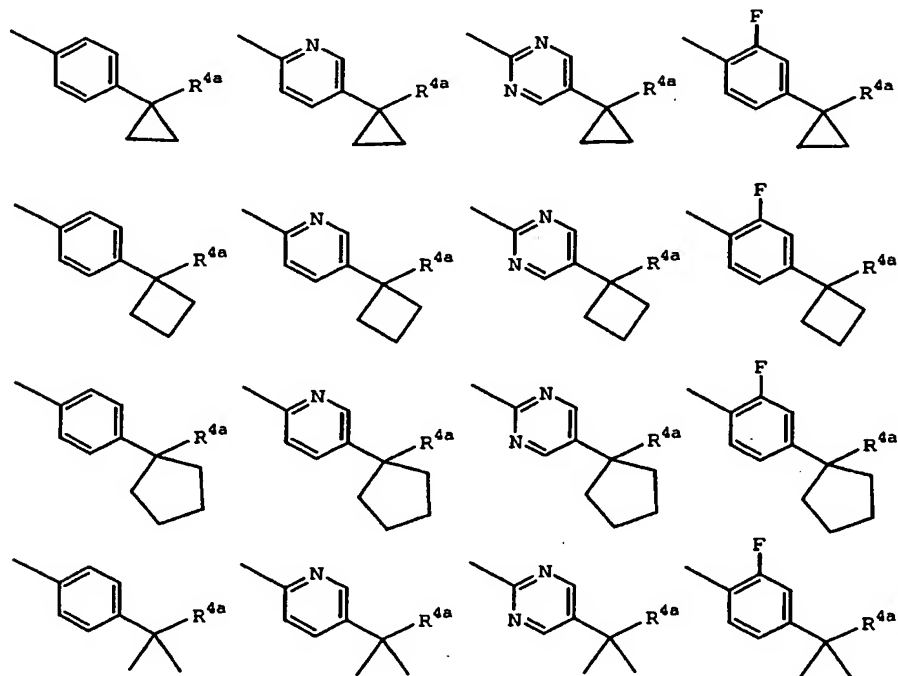
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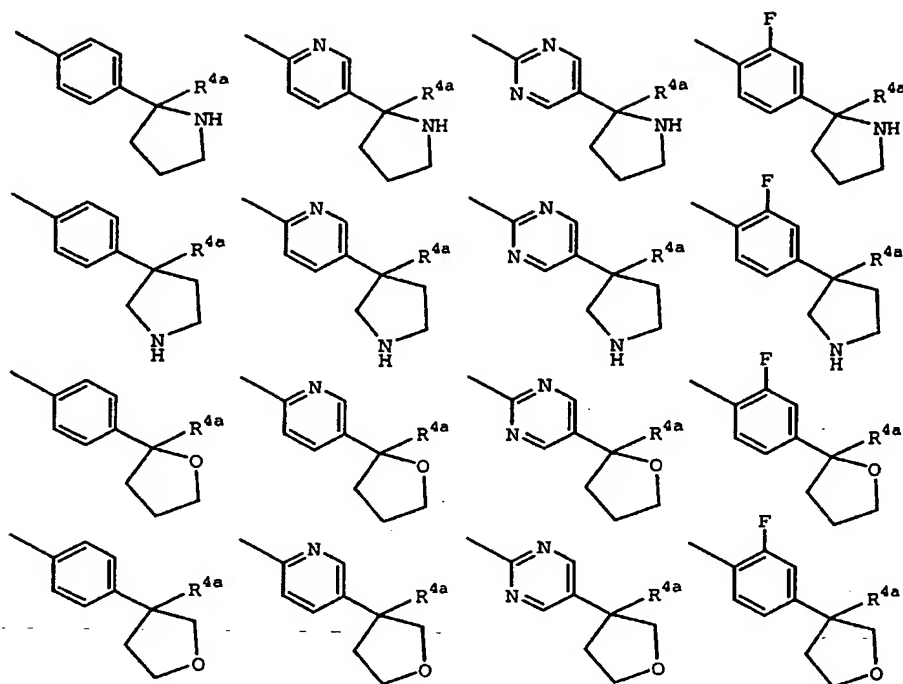


G_1 is absent or is selected from $C(O)NH$, $NHC(O)$, and $NHSO_2$;

5

A-B is selected from:





Z is selected from a bond, CH₂, and CH₂CH₂;

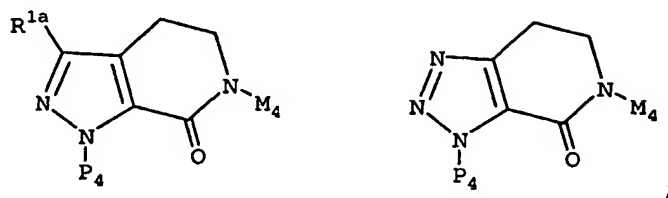
- 5 R^{2d}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl substituted with 0-2 R^{4c}, and a 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2d} forms other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;
- 10
- 15 R^{2e}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted with 0-2 R^{4c}, phenyl, substituted with 0-2 R^{4c}, and 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;
- 20

R^{4a} is selected from $NR^{2d}R^{2d}$, $CH_2NR^{2d}R^{2d}$, $CH_2CH_2NR^{2d}R^{2d}$,
 $N(\rightarrow O)R^{2d}R^{2d}$, $CH_2N(\rightarrow O)R^{2d}R^{2d}$, CH_2OR^{2d} , $C(O)R^{2e}$,
 $C(O)NR^{2d}R^{2d}$, $CH_2C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)R^{2e}$, $CH_2NR^{2d}C(O)R^{2e}$,
5 $NR^{2d}C(O)NR^{2d}R^{2d}$, $CH_2NR^{2d}C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)OR^{2d}$,
 $CH_2NR^{2d}C(O)OR^{2d}$, $NR^{2d}SO_2R^{2d}$, $CH_2NR^{2d}SO_2R^{2d}$, $S(O)_pR^{2d}$,
 $CH_2S(O)_pR^{2d}$, 5-6 membered carbocycle substituted with
0-2 R^{4c} , $-(CH_2)$ -5-6 membered carbocycle substituted
with 0-2 R^{4c} , $-(CH_2)_2$ -5-6 membered carbocycle
10 substituted with 0-2 R^{4c} , 5-6 membered heterocycle
substituted with 0-2 R^{4c} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and $S(O)_p$, $-(CH_2)$ -5-6 membered
heterocycle substituted with 0-2 R^{4c} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the
group consisting of N, O, and $S(O)_p$, and $-(CH_2)_2$ -5-6
membered heterocycle substituted with 0-2 R^{4c} and
consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and $S(O)_p$
20 provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or
 $S(O)H$; and,

R^{4c} is selected from $=O$, OH , OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$,
 $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH=CH_2$,
25 $CH\equiv CH$, CH_2OH , CH_2OCH_3 , $CH_2OCH_2CH_3$, $CH_2OCH_2CH_2CH_3$,
 $CH_2OCH(CH_3)_2$, F , Br , Cl , CF_3 , $NR^{2a}R^{2a}$, $CH_2NR^{2a}R^{2a}$,
 $C(O)R^{2c}$, $CH_2C(O)R^{2c}$, $NR^{2c}C(O)R^{2b}$, $CH_2NR^{2c}C(O)R^{2b}$,
 $C(O)NR^{2a}R^{2a}$, $CH_2C(O)NR^{2a}R^{2a}$, $SO_2NR^{2a}R^{2a}$, $CH_2SO_2NR^{2a}R^{2a}$,
 $NR^{2a}SO_2R^{5a}$, $CH_2NR^{2a}SO_2R^{5a}$, $S(O)_pR^{5a}$, and $CH_2S(O)_pR^{5a}$.

30

7. A compound according to Claim 6, wherein the compound is selected from:

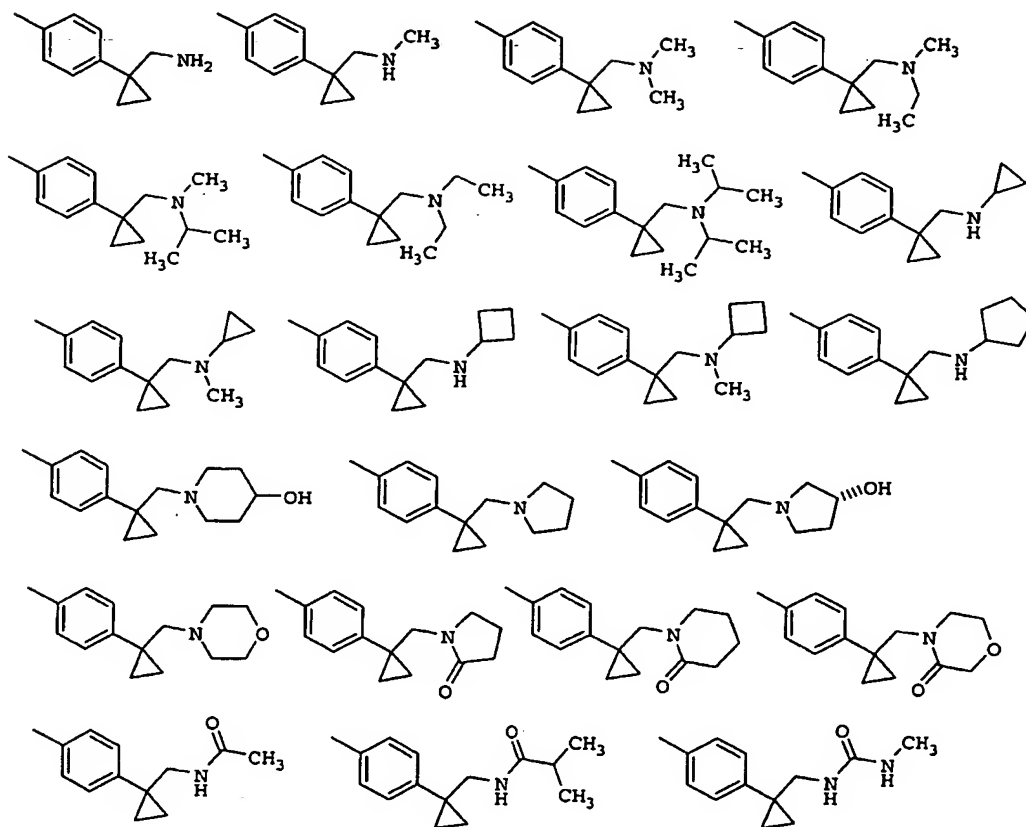


P_4 is -G;

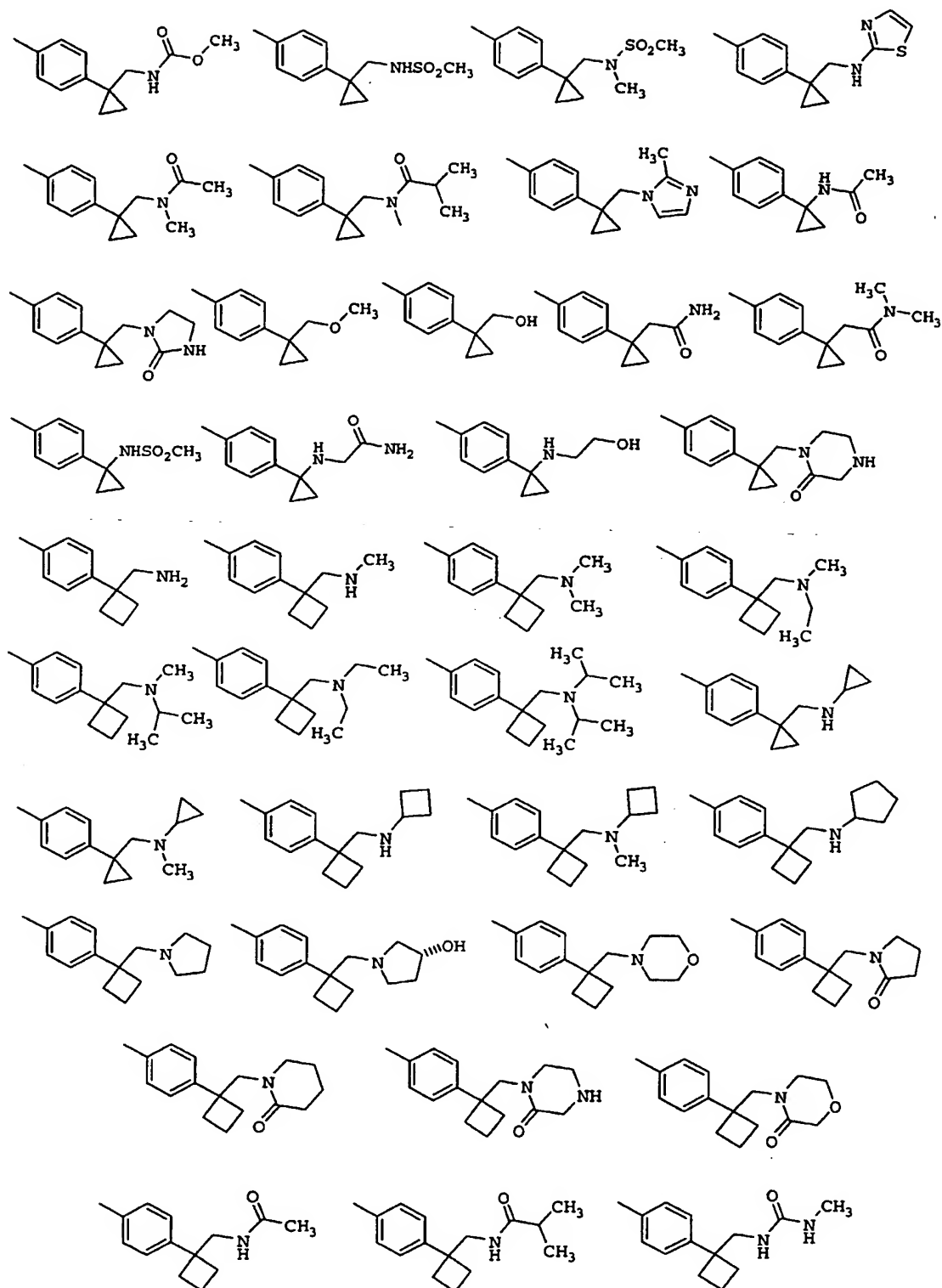
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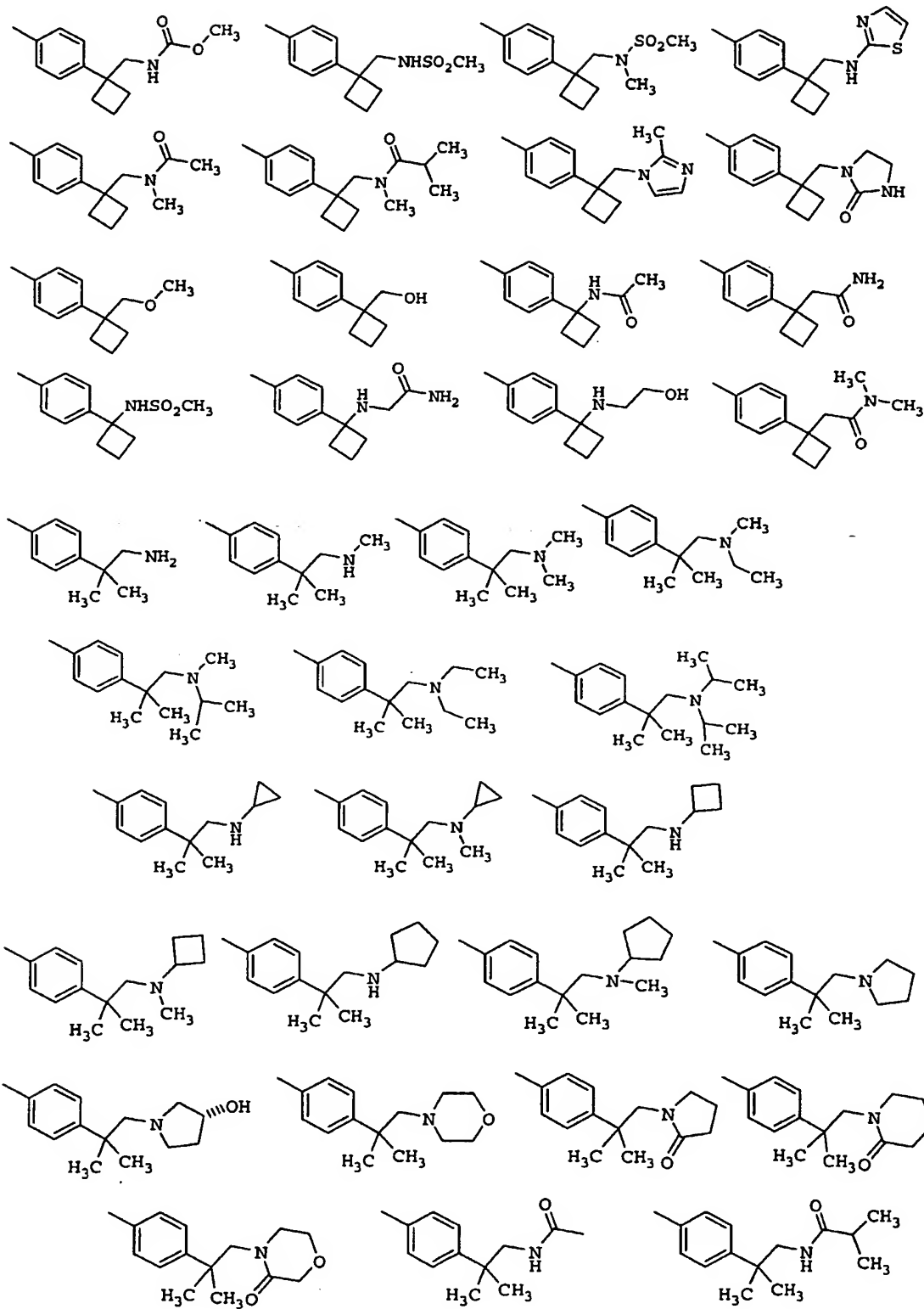
M_4 is -A-B;

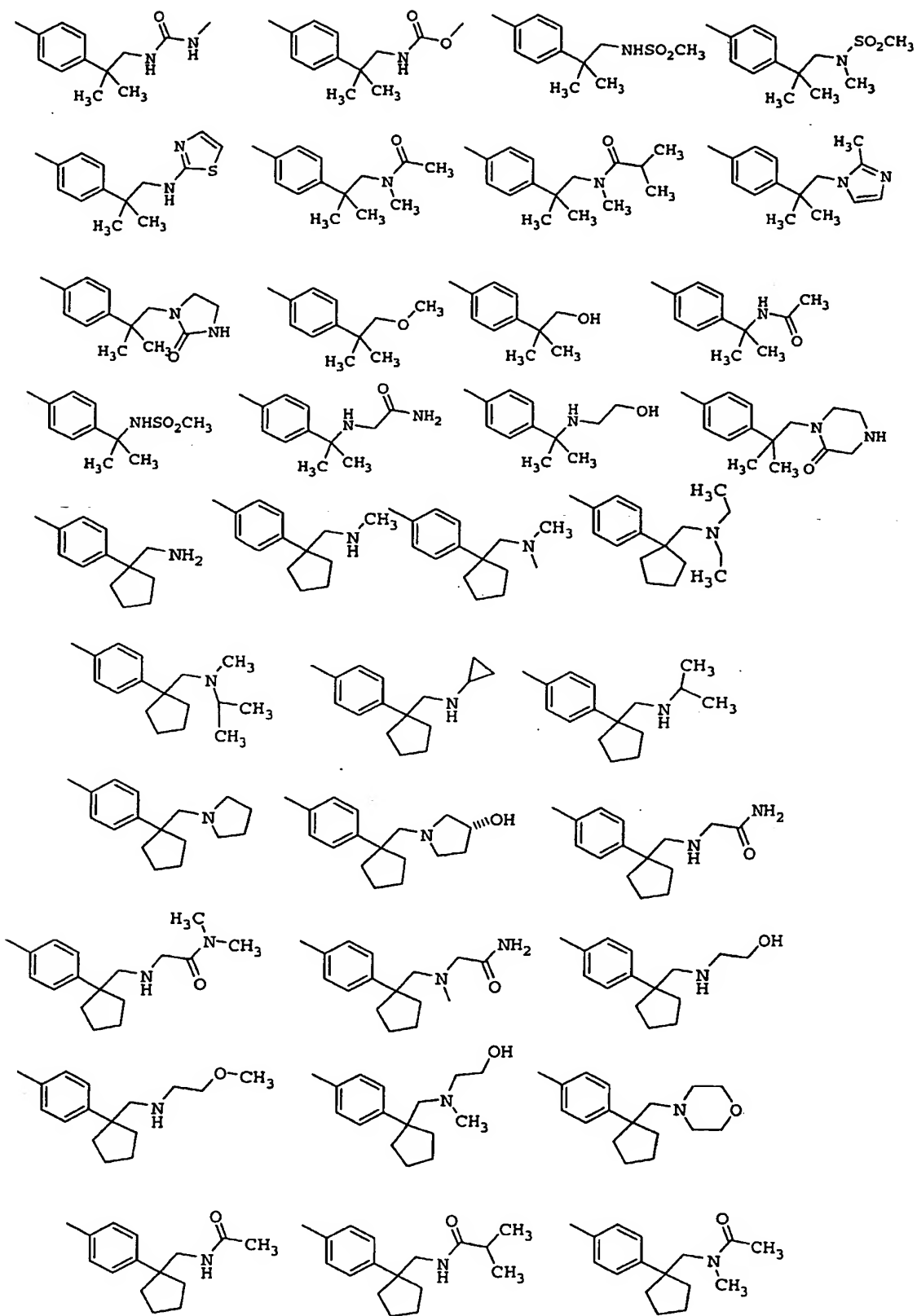
A-B is selected from:

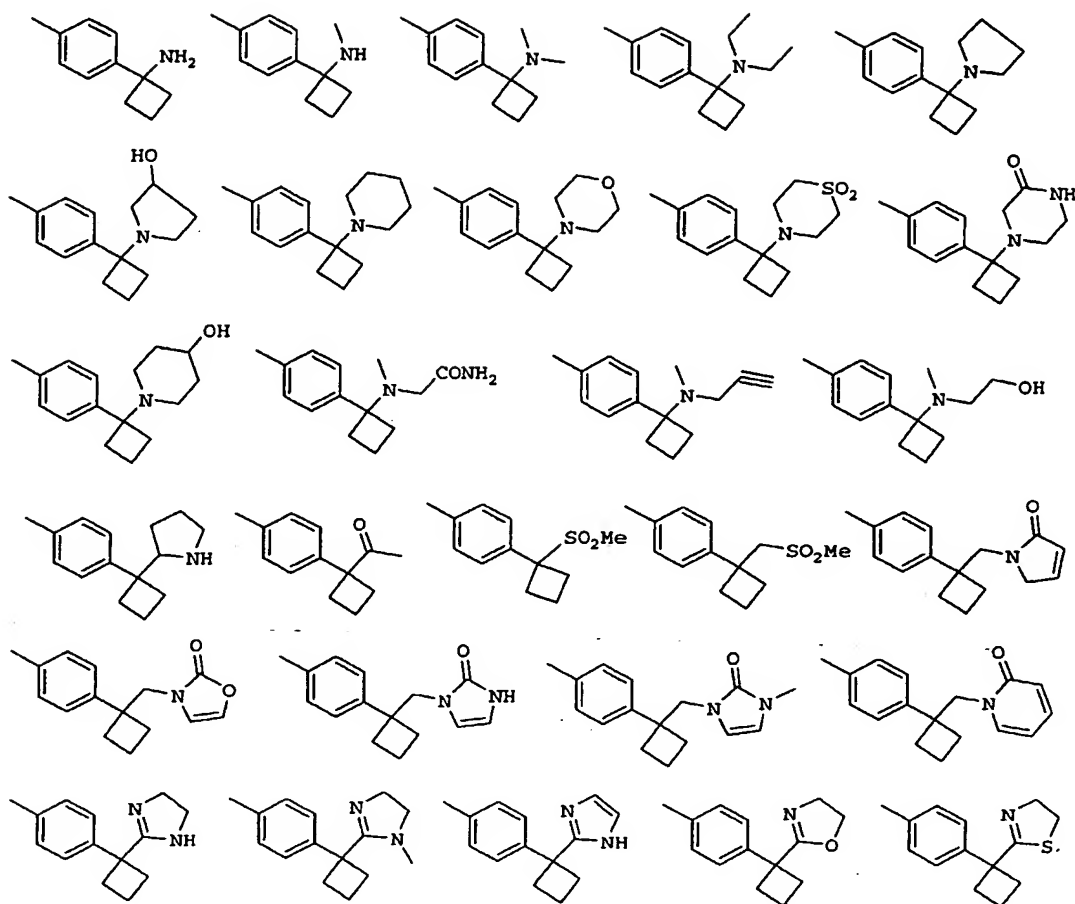


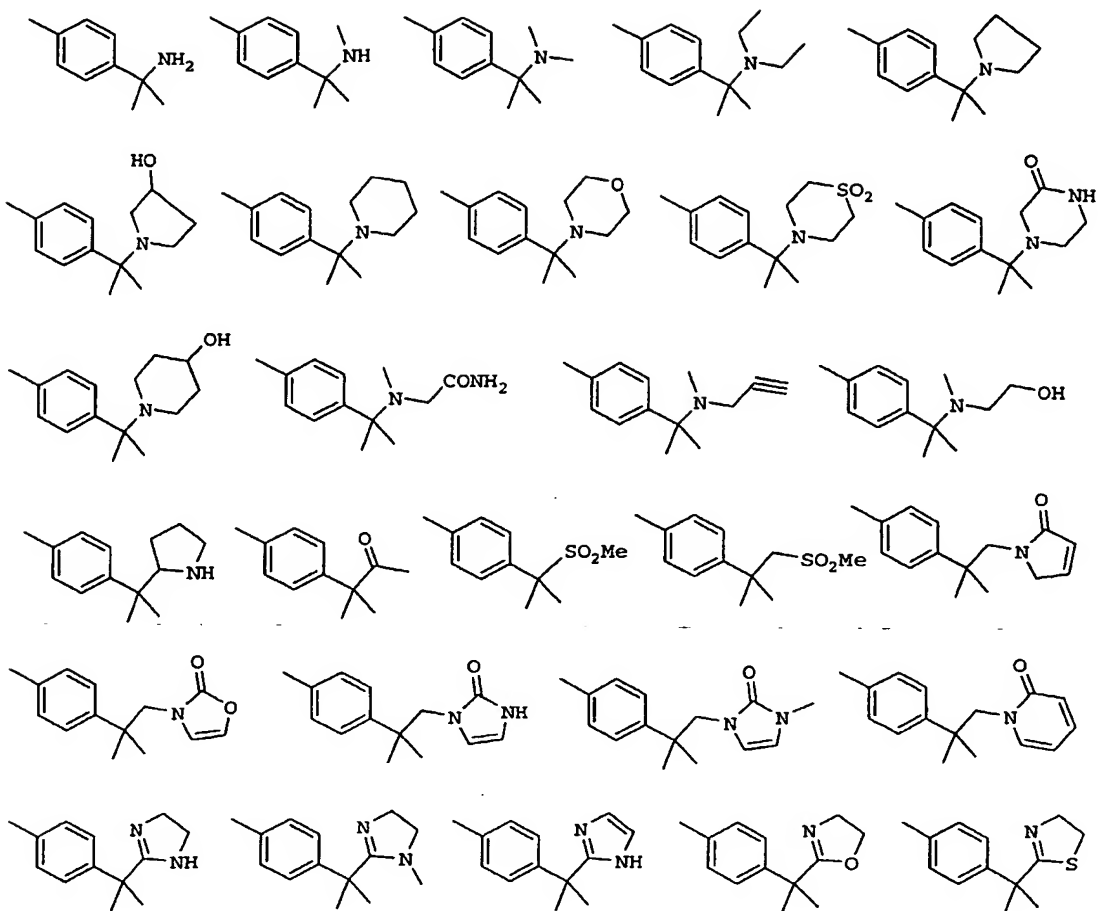
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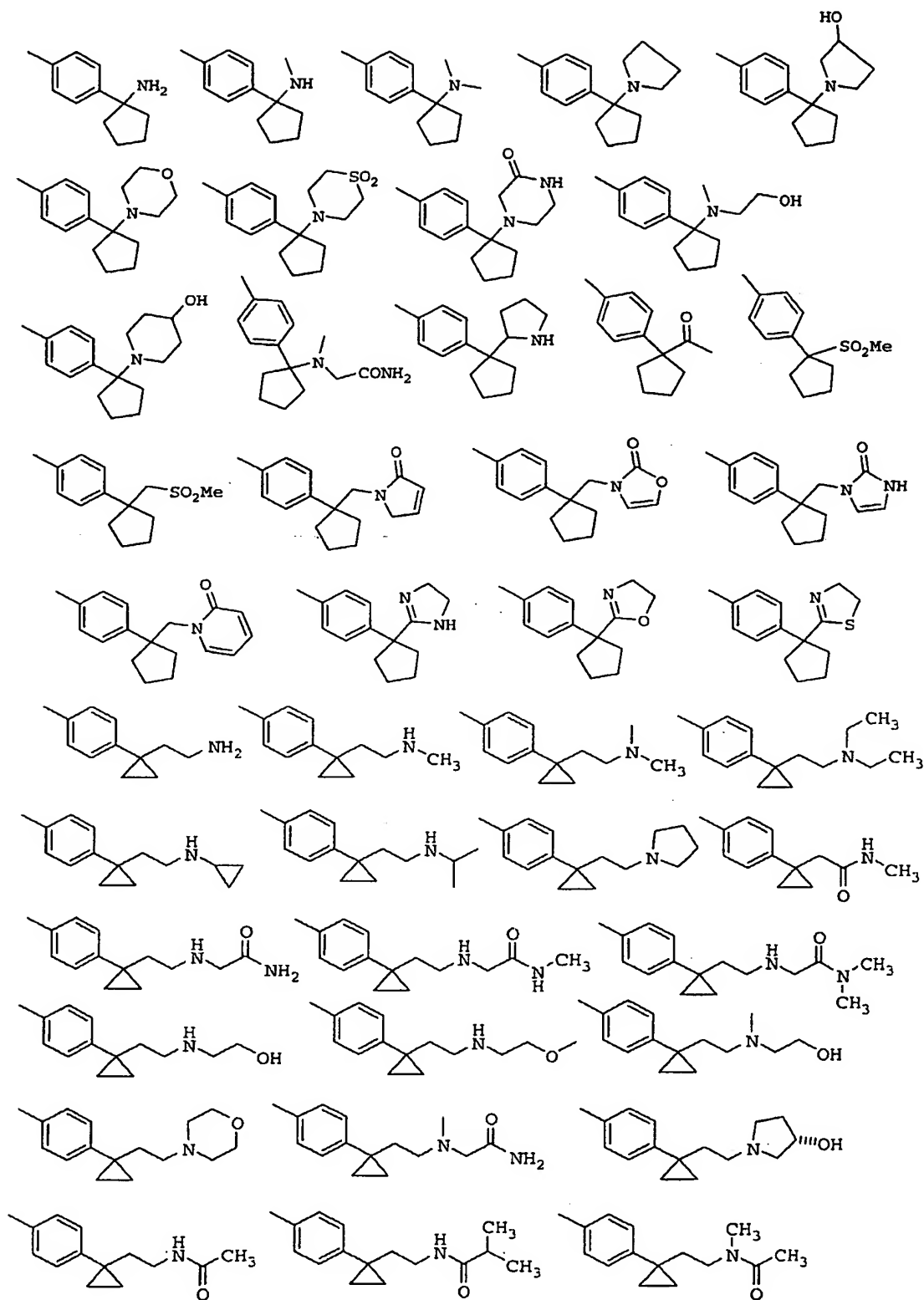


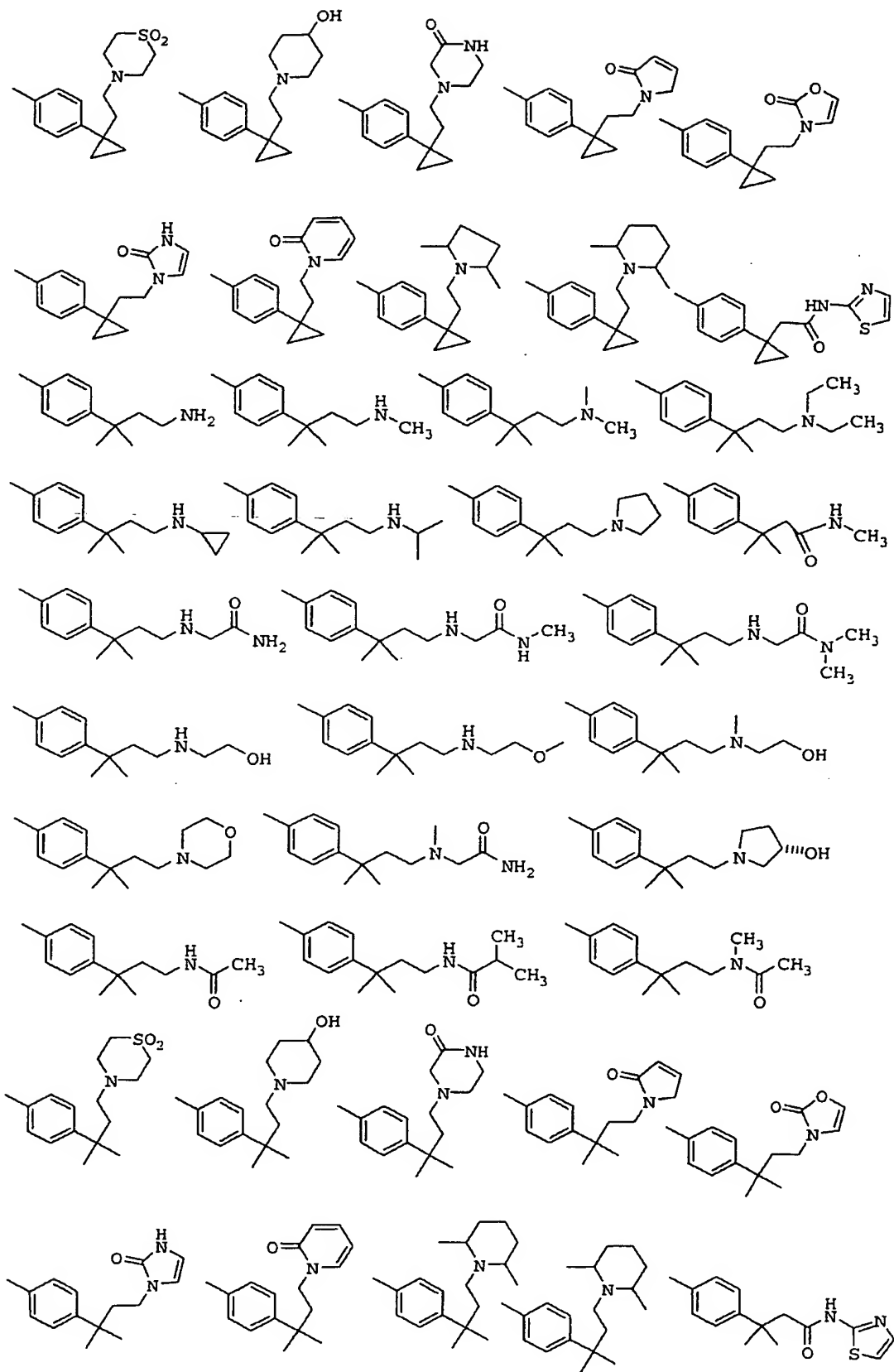


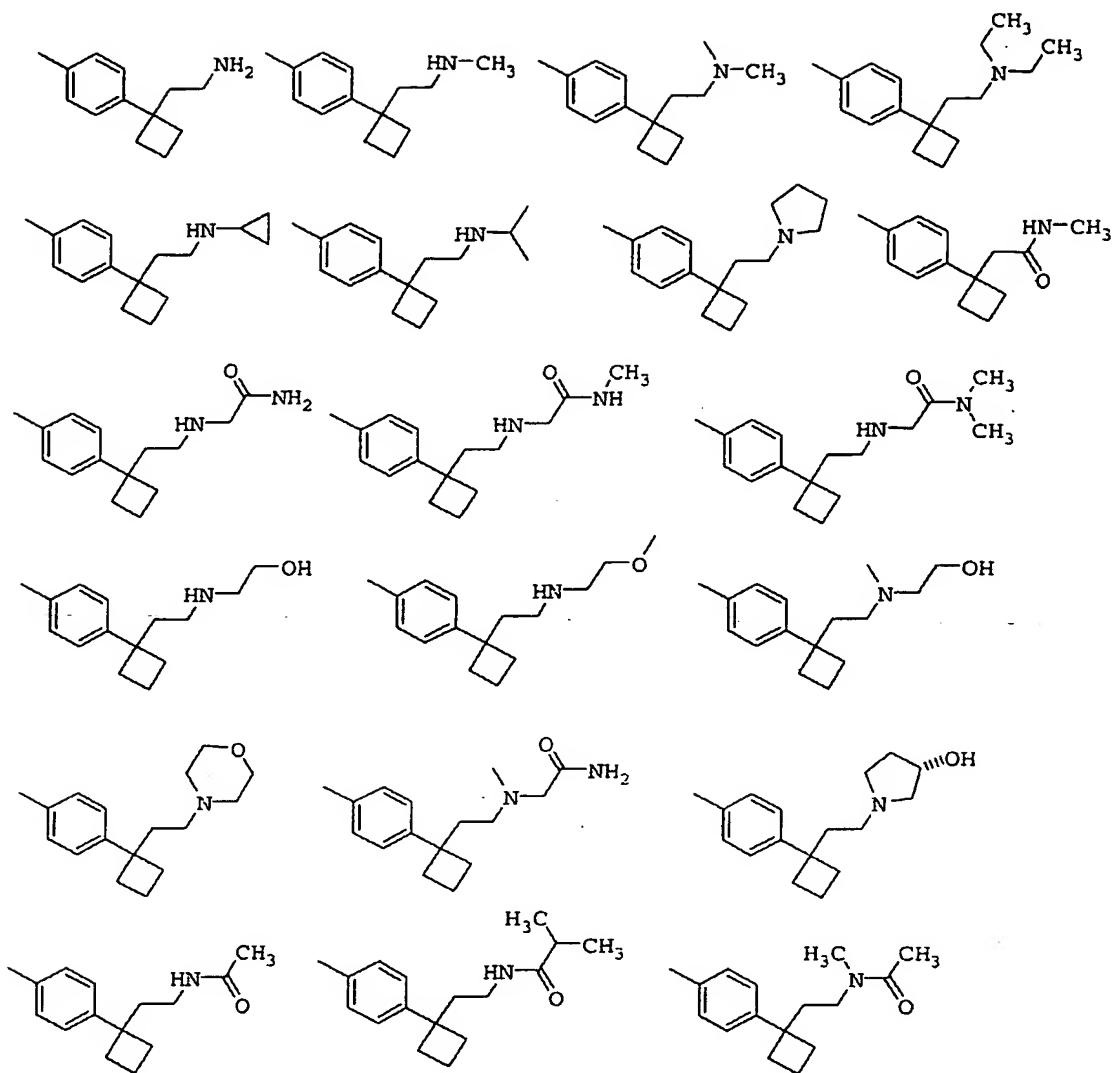


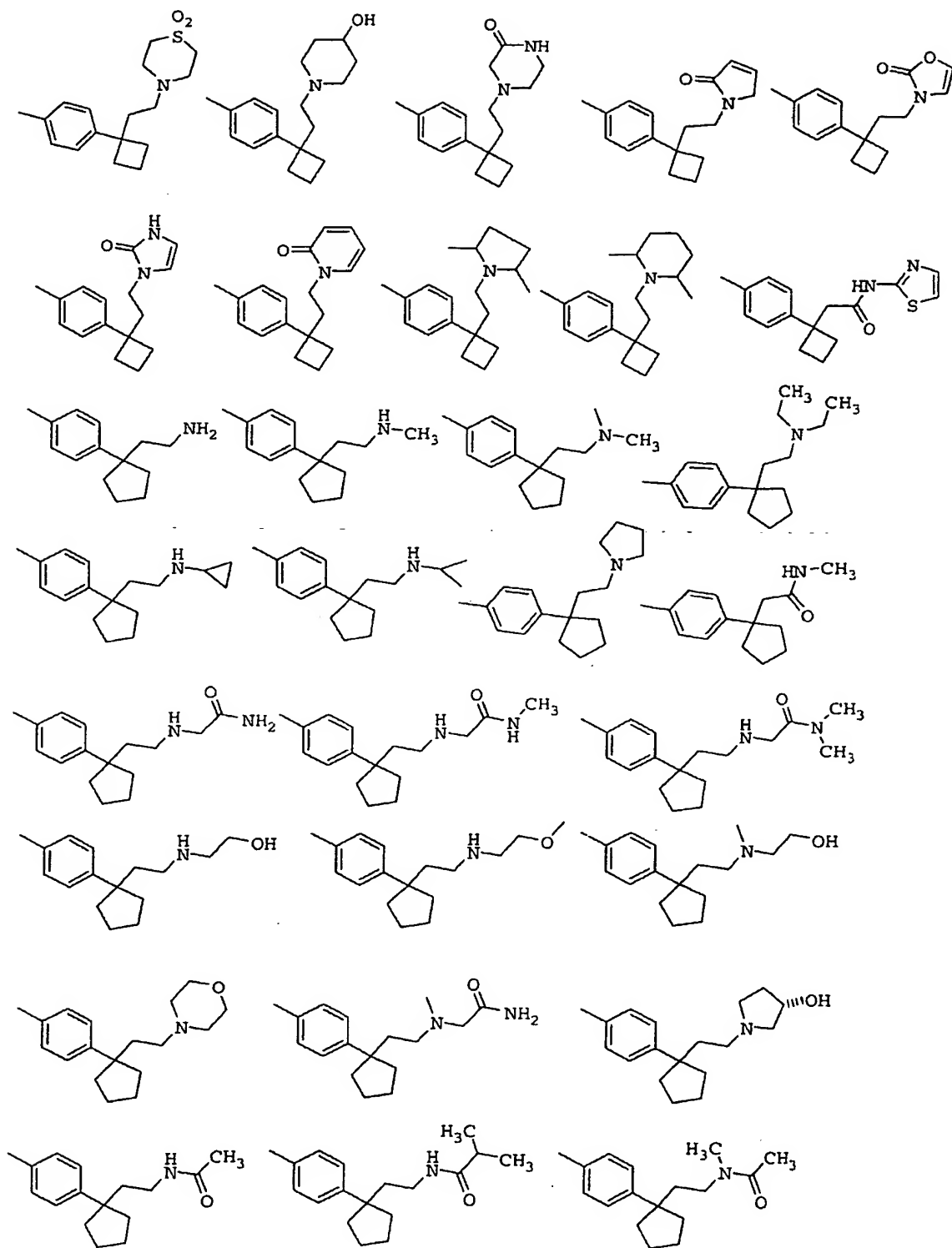


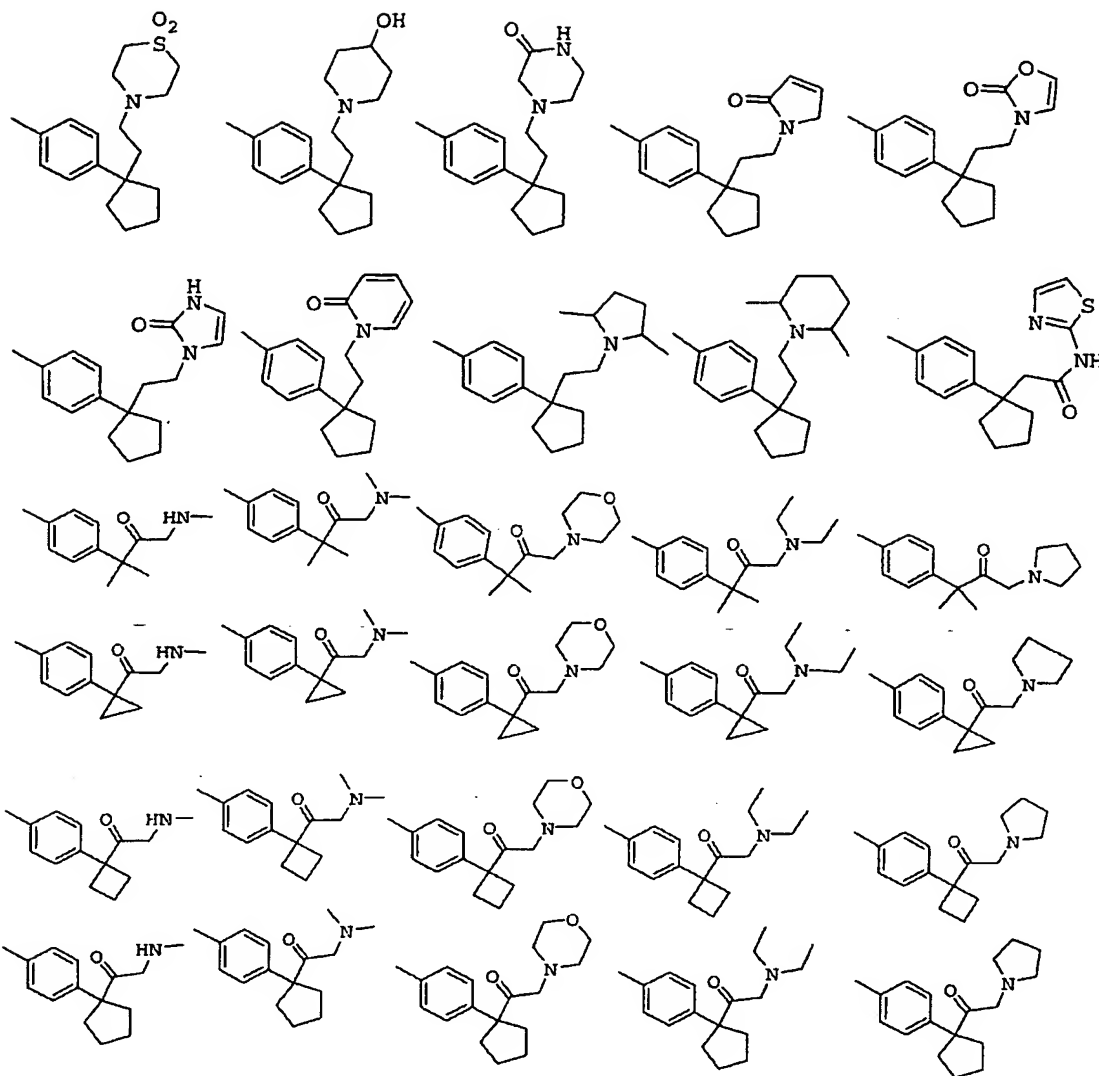












5

8. A compound according to Claim 1, wherein the compound is the compound is selected from the group:

10 1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopropyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

- 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-(4-{1-[(cyclopentylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-(4-{1-[(4-hydroxy-1-piperidinyl)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 35 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

- 1-(4-methoxyphenyl)-6-{4-[1-(1-pyrrolidinylcarbonyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-N,N-dimethylcyclopropanecarboxamide;
- 10 1-(4-methoxyphenyl)-6-(4-{1-[(4-methyl-1-piperazinyl)carbonyl]cyclopropyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(4-hydroxypiperidine-1-carbonyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropanecarboxamide;
- 25 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropanecarboxylic acid
cyclopentylamide;
- 30 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-N-(1,3,4-thiadiazol-2-yl)cyclopropanecarboxamide;
- 35 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-

yl]phenyl}-N-(1H-tetraazol-5-
yl)cyclopropanecarboxamide;

5 methyl 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropanecarboxylate;

10 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropanecarbonitrile;

15 6-{4-[1-(aminomethyl)cyclopropyl]phenyl}-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)methyl]-N-methylacetamide;

20 N'-ethyl-N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-
(trifluoromethyl)-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-
c]pyridin-6-yl]phenyl}cyclopropyl)methyl]-N-
methylurea;

25 N-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)methyl]-N-
methylethanesulfonamide;

30 1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-
ylmethyl)cyclopropyl]phenyl}-3-trifluoromethyl-
1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

35 1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)-
cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-
tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

methyl 1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl)cyclopentanecarboxylate;

5

1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclopentyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

10

6-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

15

1-(4-methoxyphenyl)-6-{4-[1-(1-
pyrrolidinylmethyl)cyclopentyl]phenyl}-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

20

6-[4-(1-[(3R)-3-hydroxy-1-
pyrrolidinyl]methyl)cyclopentyl]phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

25

1-(4-methoxyphenyl)-6-{4-[1-(4-
morpholinylmethyl)cyclopentyl]phenyl}-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

30

N-[(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl]phenyl)cyclopentyl)methyl]-N-methylacetamide;

35

N-[(1-(4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-

yl}phenyl)cyclopentyl)methyl]-N-
methylnmethanesulfonamide;

5 methyl 1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-
1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-
yl}phenyl)cyclobutanecarboxylate;

10 1-(4-methoxyphenyl)-6-(4-{1-
[(methylamino)methyl]cyclobutyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

15 6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

20 6-(4-{1-[(diethylamino)methyl]cyclobutyl}phenyl)-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

25 1-(4-methoxyphenyl)-6-{4-[1-(1-
pyrrolidinyl)methyl]cyclobutyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

30 6-[4-(1-{[(3R)-3-hydroxy-1-
pyrrolidinyl]methyl}cyclobutyl)phenyl]-1-(4-
methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-
7H-pyrazolo[3,4-c]pyridin-7-one;

35 1-(4-methoxyphenyl)-6-{4-[1-(4-
morpholinyl)methyl]cyclobutyl}phenyl)-3-
(trifluoromethyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-
c]pyridin-7-one;

- N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclobutyl)methyl]-*N*-methylacetamide;
- 5 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclobutyl)methyl]-*N*-methylmethanesulfonamide;
- 10 1-{4-[1-(4-Methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]-phenyl}cyclohexanecarboxylic acid methyl ester;
- 15 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclohexyl}phenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;
- 20 6-(4-{1-[(dimethylamino)methyl]cyclohexyl}phenyl)-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;
- 25 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclohexyl)methyl]-*N*-methylacetamide;
- 30 *N*-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-(trifluoromethyl)-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclohexyl)methyl]-*N*-methylmethanesulfonamide;
- 35 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7*H*-pyrazolo[3,4-*c*]pyridin-7-one;

- 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 5 6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-(4-{1-[(cyclopentylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;
- 30 N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)methyl]-N-methylacetamide;
- 35 3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(2-methylimidazol-1-ylmethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

3-methanesulfonyl-1-(4-methoxyphenyl)-6-{4-[1-(thiazol-2-ylaminomethyl)cyclopropyl]phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

5

1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclobutyl}phenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro 7H-pyrazolo[3,4-c]pyridin-7-one;

10

6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

15

6-(4-{1-[(isopropylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

20

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclobutyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

25

6-[4-(1-{[(3R)-3-hydroxy-1-pyrrolidinyl]methyl}cyclobutyl)phenyl]-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

30

6-(4-{1-[(diethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

35

1-(4-methoxyphenyl)-3-(methylsulfonyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-1,4,5,6-tetrahydro-7H-pyrazolo[3,4-c]pyridin-7-one;

N-[(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6*H*-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)methyl]-*N*-methylacetamide;

5 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

10 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

15 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

20

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

25

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

30 6-[4-(1-{[(3*R*)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

35 1-(4-methoxyphenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-

4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

5 6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

10 6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

15 1-(4-methoxyphenyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

20 6-[4-(1-{[(3*R*)-3-hydroxy-1-pyrrolidinyl]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

25 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

30 1-(3-chlorophenyl)-6-{4-[1-(isopropylamino)methyl]cyclopropyl}phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

35 1-(3-chlorophenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

6-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

5

1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

10

1-(3-chlorophenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

15

1-(3-chlorophenyl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

20

1-(3-chlorophenyl)-6-(4-{1-[(diethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

25

1-(3-chlorophenyl)-6-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

30

1-(3-chlorophenyl)-6-(4-{1-[(cyclopropylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

35

1-(3-chlorophenyl)-6-(4-{1-[(cyclobutylamino)methyl]cyclopropyl}phenyl)-7-oxo-

4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-6-(4-[1-(methoxymethyl)cyclopropyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;

1-(3-chlorophenyl)-6-(4-[1-(methoxymethyl)cyclopropyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

1-(3-chlorophenyl)-6-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

1-(3-chlorophenyl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carbonitrile;

1-(3-chlorophenyl)-6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-

7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

5 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

10 N-[(1-(4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)cyclopropyl)methyl]-N-methylacetamide;

15 6-(4-{1-[(cyclopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

6-(4-{1-[(cyclobutylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

20 6-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

25 6-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

30 6-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

35 5-(4-{1-[(diisopropylamino)methyl]cyclopropyl}phenyl)-3-(4-methoxyphenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

- 5-(4-{1-[(isopropylamino)methyl]cyclopropyl}phenyl)-3-(4-methoxyphenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 5 3-(4-methoxyphenyl)-5-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 10 3-(4-methoxyphenyl)-5-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 15 3-(4-methoxyphenyl)-5-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 20 5-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-3-(4-methoxyphenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 3-(3-chlorophenyl)-5-[4-(1-{[(2-hydroxyethyl)amino]methyl}cyclopropyl)phenyl]-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 25 3-(3-chlorophenyl)-5-[4-(1-{[(2-hydroxyethyl)(methyl)amino]methyl}cyclopropyl)phenyl]-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 30 3-(3-chlorophenyl)-5-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 35 3-(3-chlorophenyl)-5-(4-{1-[(3-hydroxy-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-3,5,6,7-tetrahydro-4H-[1,2,3]triazolo[4,5-c]pyridin-4-one;

- 6-[4-(1-[(2-hydroxyethyl)(methyl)amino]methyl)cyclopropyl]phenyl]-
1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
5 pyrazolo[3,4-c]pyridine-3-carboxamide;
- 6-{4-[1-(dimethylamino)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxamide;
10
- 6-(4-{1-[(2-hydroxyethyl)(methyl)amino]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 2-(1-{4-[1-(4-methoxyphenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
20 pyrazolo[3,4-c]pyridine-3-carboxamide;
- 2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
25
- 2-(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)acetamide;
30
- 2-(1-{4-[1-(3-chlorophenyl)-3-cyano-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)acetamide;

- 1-(3-chlorophenyl)-6-(4-{1-[2-(dimethylamino)-2-oxoethyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 2-(1-{4-[1-(3-chlorophenyl)-3-(methylsulfonyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 10 2-(1-{4-[3-(3-chlorophenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 15 2-(1-{4-[3-(4-methoxyphenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)-N,N-dimethylacetamide;
- 20 2-(1-{4-[3-(4-methoxyphenyl)-4-oxo-3,4,6,7-tetrahydro-5H-[1,2,3]triazolo[4,5-c]pyridin-5-yl]phenyl}cyclopropyl)acetamide;
- 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-imidazolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-piperazinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(3-oxo-4-morpholinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 35 1-(4-methoxyphenyl)-7-oxo-6-(4-{1-[(2-oxo-1-piperidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 6-(4-[1,1-dimethyl-2-(2-oxodihydro-2H-1,3-oxazin-3(4H)-yl)ethyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 1-(4-[3-(aminocarbonyl)-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-1-methylethyl methylcarbamate;
- 10 1-(4-[3-(aminocarbonyl)-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-tetrahydro-6H-pyrazolo[3,4-c]pyridin-6-yl]phenyl)-1-methylethyl 3-pyrrolidinylcarbamate;
- 15 6-(4-[1-ethyl-1-(1-pyrrolidinylmethyl)propyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 6-(4-[1-[(dimethylamino)methyl]-1-ethylpropyl]phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 1-[3-(aminomethyl)phenyl]-6-(4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 25 1-[3-(aminomethyl)phenyl]-6-(4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 30 1-[3-(aminocarbonyl)phenyl]-6-(4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 35 1-[3-(aminocarbonyl)phenyl]-6-(4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

- 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 5 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;
- 10 1-(1-amino-7-isoquinolinyl)-6-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 15 1-(1-amino-7-isoquinolinyl)-6-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 20 1-(1-amino-7-isoquinolinyl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 25 1-(1-amino-7-isoquinolinyl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 30 1-(3-amino-1,2-benzisoxazol-5-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1*H*-pyrazolo[3,4-*c*]pyridine-3-carboxamide;
- 1-(3-amino-1,2-benzisoxazol-5-yl)-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-3-(trifluoromethyl)-1,4,5,6-tetrahydro-7*H*-pyrazolo[3,4-*c*]pyridin-7-one;

- 1-(3-amino-1,2-benzisoxazol-5-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-[3-(aminomethyl)phenyl]-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 10 6-[4-(1-{[acetyl(methyl)amino]methyl}cyclopropyl)phenyl]-1-[3-(aminomethyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 1-[3-(aminocarbonyl)phenyl]-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 3-[3-cyano-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridin-1-yl]benzamide;
- 25 1-(2,3-dihydro-1H-indol-6-yl)-6-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 30 1-(2,3-dihydro-1H-indol-6-yl)-7-oxo-6-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 1-(2,3-dihydro-1H-indol-6-yl)-7-oxo-6-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;

- 6-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 5 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclobutyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 10 1-(4-methoxyphenyl)-6-{4-[1-(4-morpholinylmethyl)cyclopentyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 15 6-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxamide;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(2-oxo-pyrrolidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 1-(4-methoxyphenyl)-6-{4-[1-(2-oxo-piperidin-1-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)-phenyl]-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 N-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-cyclopropyl)-N-methyl-acetamide;

N-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-methanesulfonamide;

5 *N*-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-2-methylaminoacetamide;

10 2-dimethylamino-*N*-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}cyclopropyl)-*N*-methylacetamide;

N-(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-*c*]pyridin-6-yl]phenyl}-cyclopropyl)-*N*-methyl-2-morpholin-4-yl-acetamide;

15 6-{4-[1-(1-hydroxyethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

20 6-[4-(1-acetylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

25 6-{4-[1-(1-hydroxy-1-methyl-ethyl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

30 6-[4-(1-methoxymethylcyclopropyl)phenyl]-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-*c*]pyridin-7-one;

- 6-{4-[1-(4,5-dihydro-oxazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-(4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropanecarboxylic acid 2-amino-ethyl ester ;
- 10 6-{4-[1-(4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 6-{4-[1-(4,5-dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxyphenyl)-6-{4-[1-(1-methyl-4,5-dihydro-1H-imidazol-2-yl)cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 6-{4-[1-(1-methanesulfonyl-4,5-dihydro-1H-imidazol-2-yl)-cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-{4-[1-(1H-imidazol-2-yl)cyclopropyl]phenyl}-1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 1-(4-methoxyphenyl)-6-{4-[1-(1-methyl-1H-imidazol-2-yl)-cyclopropyl]phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 2-[(1-{4-[1-(4-methoxyphenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]phenyl}-cyclopropyl)-methyl-amino]-acetamide;

6-(4-{1-[(2-hydroxyethyl)-methylamino]cyclopropyl}phenyl)-
1-(4-methoxyphenyl)-3-trifluoromethyl-1,4,5,6-
tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

5

1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopropanecarboxylic acid methoxy-methyl-amide;

10 6-[4-(1-hydroxymethylcyclopropyl)phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amide;

15 6-[4-(1-acetyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-
oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide ;

20 6-[4-(1-aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide;

25 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)-
phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amid;

6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

30 6-[4-(1-methylaminomethylcyclopentyl)phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4c]pyridine-3-carboxylic acid amide;

35 6-[4-(1-dimethylaminomethylcyclopentyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

6-[4-(1-dimethylaminomethylcyclopentyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

5

6-[4-(1-[(2-hydroxyethyl)methylaminomethyl]-cyclopentyl)phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

10

6-[4-(1-hydroxymethyl-cyclopentyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15 6-(4-{1-[(2-hydroxyethyl)methylamino]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

20 1-(4-methoxyphenyl)-6-{4-[1-(methyl-prop-2-ynylamino)-cyclopropyl]phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

25 3-(1-hydroxyethyl)-1-(4-methoxyphenyl)-6-[4-(1-methylamino-cyclopropyl)phenyl]-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

3-acetyl-1-(4-methoxyphenyl)-6-[4-(1-methylamino-cyclopropyl)phenyl]-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

30

1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid methylamide;

- 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid dimethylamide;
- 5 6-[4-(1-aminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carbonitrile;
- 10 1-(4-methoxyphenyl)-6-[4-(1-methylaminocyclopropyl)phenyl]-
7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carbonitrile;
- 15 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-
methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 20 2-[(1-{4-[3-cyano-1-(4-methoxyphenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-
yl]phenyl}cyclopropyl)-methylamino]acetamide;
- 6-(4-{1-[(2-hydroxyethyl)methylamino]cyclopropyl}phenyl)-1-
(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-
cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 30 1-(4-methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-
cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 35 1-(4-methoxyphenyl)-7-oxo-6-[4-(1-pyrrolidin-1-yl-
cyclopropyl)phenyl]-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 1-(4-methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 5 1-(4-methoxyphenyl)-6-[4-(1-morpholin-4-yl-cyclopropyl)phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid methylamide;
- 15 6-[4-(1-dimethylaminocyclopropyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid dimethylamide;
- 20 6-[4-[1-(1,1-dioxo-1 λ ⁶-thiomorpholin-4-yl)cyclopropyl]phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-aminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 6-[4-(1-dimethylaminocyclopropylmethyl)phenyl]-1-(4-methoxyphenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(3-chloro-phenyl)-6-[4-[1,1-dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

- 6-{4-[1,1-dimethyl-2-(2-oxo-pyrrolidin-1-yl)-ethyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-(4-methoxy-phenyl)-6-[4-(1-methyl-1-pyrrolidin-1-ylethyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 10 6-[4-(1-dimethylamino-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 15 6-{4-[1-(4,4-dimethyl-4,5-dihydro-oxazol-2-yl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-[4-(1-methanesulfonyl-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-hydroxy-1-methyl-ethyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 35 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;

- 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 5 1-(4-methoxy-phenyl)-6-{4-[1-(2-oxo-2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-acetyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;

6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

5

1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;

10

1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15

6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;

20

6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

25

6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

30

1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

35

1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

35

1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

5

6-{4-[1-(2-diethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

10 1-(4-methoxy-phenyl)-7-oxo-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15 6-(4-{1-[2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

20 6-(4-{1-[2-(3-hydroxy-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

25 6-(4-{1-[2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

30 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-piperidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

35 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

1- (4-methoxy-phenyl) -6- (4- (1- [2- (methyl-thiazol-2-yl)-
amino) -ethyl] -cyclopropyl) -phenyl) -7-oxo-4,5,6,7-
tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic
acid amide;

5

6- [4- (1- {2- [(2-hydroxy-ethyl) -methyl-amino] -ethyl} -
cyclopropyl) -phenyl] -1- (4-methoxy-phenyl) -7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide;

10

1- (4-methoxy-phenyl) -6- (4- (1- {2- (2-methyl-imidazol-1-yl) -
ethyl} -cyclopropyl) -phenyl) -7-oxo-4,5,6,7-tetrahydro-
1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15

6- (4- {1- [2- (2,6-dimethyl-piperidin-1-yl) -ethyl] -
cyclopropyl} -phenyl) -1- (4-methoxy-phenyl) -7-oxo-
4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-
carboxylic acid amide;

20

2- (1- {4- [3-methanesulfonyl-1- (4-methoxy-phenyl) -7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl] -
phenyl} -cyclopropyl) -N,N-dimethyl-acetamide;

25

2- (1- {4- [3-methanesulfonyl-1- (4-methoxy-phenyl) -7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl] -
phenyl} -cyclopropyl) -acetamide;

30

2- (1- {4- [3-methanesulfonyl-1- (4-methoxy-phenyl) -7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl] -
phenyl} -cyclopropyl) -N-methyl-acetamide;

35

2- (1- {4- [3-methanesulfonyl-1- (4-methoxy-phenyl) -7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl] -
phenyl} -cyclopropyl) -N,N-dimethyl-acetamide;

- 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 3-methanesulfonyl-6-{4-[1-(2-methoxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-diethylamino-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 25 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-isopropylamino-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-{4-[1-(2-(2,5-dimethyl-pyrrolidin-1-yl)-ethyl)-cyclopropyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 6-(4-{1-[2-(3-hydroxy-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-piperidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-[4-(1-{2-[(2-hydroxy-ethyl)-methyl-amino]-ethyl}-cyclopropyl)-phenyl]-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 2-{[2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl]-cyclopropyl)-ethyl]-methyl-amino}-acetamide;
- 25 2-[2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl)-cyclopropyl)-ethylamino]-acetamide;
- 30 6-(4-{1-[2-(2-hydroxy-ethylamino)-ethyl]-cyclopropyl}-phenyl)-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-methyl-imidazol-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(thiazol-2-ylamino)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 3-methanesulfonyl-1-(4-methoxy-phenyl)-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 15 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;
- 20 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 35 1-(4-methoxy-phenyl)-7-oxo-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

- 1- (4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 6-(4-{1-[2-(1,1-dioxo-1,1,6-thiomorpholin-4-yl)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-(4-{1-[2-(2-hydroxy-ethylamino)-ethyl]-cyclopropyl}-phenyl)-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 15 2-[2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethylamino]-acetamide;
- 20 2-{[2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethyl]-methyl-amino}-acetamide;
- 25 6-[4-(1-{2-[(2-hydroxy-ethyl)-methyl-amino]-ethyl}-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 30 N-[2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethyl]-N-methyl-methanesulfonamide;
- N-[2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-ethyl]-N-methyl-acetamide;

- 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 1-(4-methoxy-phenyl)-7-oxo-6-(4-{1-[2-(2-oxo-2H-pyridin-1-yl)-ethyl]-cyclopropyl}-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 10 6-(4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl)-1-(4-methoxy-phenyl)-3-methyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-(4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl)-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 5-(4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl)-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 5-(4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl)-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;
- 25 6-(4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl)-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-(4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl)-1-(4-methoxy-phenyl)-3-methyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-(4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl)-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 5 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 10 6-{4-[1-(2-dimethylamino-ethyl)-cyclobutyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-3-methanesulfonyl-1-(4-methoxy-phenyl)-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;
- 25 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 35 6-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-1-(4-methoxy-phenyl)-3-methyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5-{4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-3-(4-methoxy-phenyl)-3,5,6,7-tetrahydro-[1,2,3]triazolo[4,5-c]pyridin-4-one;

- 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-N-methyl-acetamide;
- 5 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-N,N-dimethyl-acetamide;
- 10 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclopentyl)-acetamide;
- 15 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-
phenyl}-cyclopentyl)-acetamide;
- 20 6-[4-(1-carbamoylmethyl-cyclopentyl)-phenyl]-1-(4-methoxy-
phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-
c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-
cyclopentyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 25 6-[4-(1-dimethylcarbamoylmethyl-cyclopentyl)-phenyl]-1-(4-
methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-
pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 30 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopentyl)-N,N-dimethyl-acetamide;
- 35 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclopentyl)-N-methyl-acetamide;

- 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-N-methyl-acetamide;
- 5 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-N,N-dimethyl-acetamide;
- 10 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopentyl)-acetamide;
- 15 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;
- 20 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 25 2-(1-{4-[3-methanesulfonyl-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N,N-dimethyl-acetamide;
- 30 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N-methyl-acetamide;
- 35 2-(1-{4-[3-cyano-1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-acetamide;

- 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-
phenyl}-cyclobutyl)-acetamide;
- 5 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-
1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-
phenyl}-cyclobutyl)-N-methyl-acetamide;
- 10 2-(1-{4-[1-(4-methoxy-phenyl)-3-methyl-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclobutyl)-N-methyl-acetamide;
- 15 2-(1-{4-[1-(4-methoxy-phenyl)-3-methyl-7-oxo-1,4,5,7-
tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-
cyclobutyl)-acetamide;
- 20 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-
pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-
acetamide;
- 25 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-1,4,5,7-tetrahydro-
pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclobutyl)-N,N-
dimethyl-acetamide;
- 30 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclobutyl)-N,N-dimethyl-acetamide;
- 35 2-(1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclobutyl)-N-methyl-acetamide;

- 2- (1-{4-[3-(4-methoxy-phenyl)-4-oxo-3,4,6,7-tetrahydro-
[1,2,3]triazolo[4,5-c]pyridin-5-yl]-phenyl}-
cyclobutyl)-acetamide;
- 5 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-
dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-
2,3-dihydro-1H-isoindol-4-yl]-amide;
- 10 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-
dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-
dihydro-1H-isoindol-4-yl]-amide;
- 15 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-
dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-
dihydro-1H-isoindol-4-yl]-amide;
- 20 5-chloro-thiophene-2-carboxylic acid [2-(2-{4-[1-(2-
dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-
dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 25 5-chloro-thiophene-2-carboxylic acid [2-(2-{4-[1-(2-
dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-
oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 30 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-
dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-
dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 35 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-
dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1-
oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;

- 5-chloro-thiophene-2-carboxylic acid [2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid {2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;

- 5-chloro-thiophene-2-carboxylic acid {2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl}-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1-oxo-2,3-dihydro-1H-isoindol-4-yl}-amide;

- 5-chloro-thiophene-2-carboxylic acid {6-chloro-2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 5 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 10 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 15 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 20 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 25 5-chloro-thiophene-2-carboxylic acid [6-chloro-2-(2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-ethyl)-3-oxo-2,3-dihydro-1H-isoindol-4-yl]-amide;
- 30 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[4-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{2-[3-(1-dimethylaminomethyl-cyclopropyl)-phenyl]-ethyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 25 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-[3-(1-dimethylaminomethyl-cyclopropyl)-benzyl]-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 30 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 35 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;

- 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1,3-dioxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-1-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (6-chloro-2-{3-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzyl}-3-oxo-2,3-dihydro-1H-isoindol-4-yl)-amide;
- 20 (1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetic acid;
- 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-acetamide;
- 25 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N-methyl-acetamide;
- 30 2-(1-{4-[1-(4-methoxy-phenyl)-7-oxo-3-trifluoromethyl-1,4,5,7-tetrahydro-pyrazolo[3,4-c]pyridin-6-yl]-phenyl}-cyclopropyl)-N,N-dimethyl-acetamide;
- 35 1-(4-methoxy-phenyl)-6-{4-[1-(2-oxo-2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;

- 6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 5 1-(4-methoxy-phenyl)-6-{4-[1-(2-methylamino-ethyl)cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 10 6-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 15 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 20 1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one
- 25 1-(4-methoxy-phenyl)-6-{4-[1-(2-pyrrolidin-1-yl-acetyl)-cyclopropyl]-phenyl}-3-trifluoromethyl-1,4,5,6-tetrahydro-pyrazolo[3,4-c]pyridin-7-one;
- 30 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 35 6-[4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;
- 1-(4-methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;
- 1-(4-Methoxy-phenyl)-6-[4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid ethyl ester;

5

6-[4-(1-dimethylcarbamoylmethyl-cyclopropyl)-phenyl]-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

10

6-{4-[1-(2-hydroxy-ethyl)-cyclopropyl]-phenyl}-1-(4-methoxy-phenyl)-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide;

15

1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carboxylic acid amide; and,

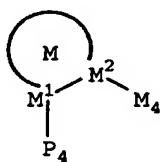
20

1-(4-methoxy-phenyl)-6-{4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-7-oxo-4,5,6,7-tetrahydro-1H-pyrazolo[3,4-c]pyridine-3-carbonitrile;

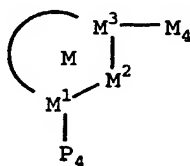
or a pharmaceutically acceptable salt form thereof.

25

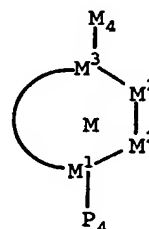
9. A compound according to Claim 1, wherein the compound is of Formula IIIa, IIIb, or IIIc:



IIIa



IIIb



IIIc

or a stereoisomer or pharmaceutically acceptable salt thereof, wherein;

30

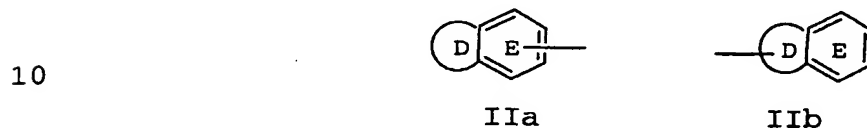
ring M, including M₁, M₂, and, if present, M₃, is phenyl or a 3-10 membered carbocyclic or 4-10 membered

heterocyclic ring consisting of: carbon atoms and 1-4 heteroatoms selected from O, S(O)_p, N, and NZ²;

ring M is substituted with 0-3 R^{1a} and 0-2 carbonyl groups,
5 and there are 0-3 ring double bonds;

one of P₄ and M₄ is -Z-A-B and the other -G₁-G;

G is a group of formula IIa or IIb:



ring D, including the two atoms of Ring E to which it is attached, is a 5-6 membered ring consisting of carbon
15 atoms and 0-2 heteroatoms selected from the group consisting of N, O, and S(O)_p;

ring D is substituted with 0-2 R and there are 0-3 ring double bonds;

20

E is selected from phenyl, pyridyl, pyrimidyl, pyrazinyl, and pyridazinyl, and is substituted with 1-3 R;

alternatively, ring D is absent, and ring E is selected
25 from phenyl, pyridyl, pyrimidyl, and thienyl, and ring E is substituted with 1-3 R;

alternatively, ring D is absent, ring E is selected from phenyl, pyridyl, and thienyl, and ring E is
30 substituted with 1 R and substituted with a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p, wherein the 5-6 membered heterocycle

is substituted with 0-2 carbonyls and 1-3 R and there are 0-3 ring double bonds;

R is selected from H, C₁₋₄ alkyl, F, Cl, OH, OCH₃, OCH₂CH₃,
 5 OCH(CH₃)₂, CN, C(=NH)NH₂, C(=NH)NHOH, C(=NH)NHOCH₃,
 NH₂, NH(C₁₋₃ alkyl), N(C₁₋₃ alkyl)₂, C(=NH)NH₂, CH₂NH₂,
 CH₂NH(C₁₋₃ alkyl), CH₂N(C₁₋₃ alkyl)₂, (CR⁸R⁹)_tNR⁷R⁸,
 C(O)NR⁷R⁸, CH₂C(O)NR⁷R⁸, S(O)_pNR⁷R⁸, CH₂S(O)_pNR⁷R⁸,
 SO₂R³, and OCF₃;

10

alternatively, when 2 R groups are attached to adjacent atoms, they combine to form methylenedioxy or ethylenedioxy;

15 A is selected from:

C₅₋₁₀ carbocycle substituted with 0-2 R⁴, and

5-10 membered heterocycle substituted with 0-2 R⁴ and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

20

X is selected from -(CR²R^{2a})₁₋₄-, -C(O)-, -C(O)CR²R^{2a}-,
 -CR²R^{2a}C(O), -S(O)₂-, -S(O)₂CR²R^{2a}-, -CR²R^{2a}S(O)₂-,
 -NR²S(O)₂-, -S(O)₂NR²-, -NR²C(O)-, -C(O)NR²-, NR²,
 -NR²CR²R^{2a}-, -CR²R^{2a}NR²-, O, -OCR²R^{2a}-, and -CR²R^{2a}O-;

25

Y is a C₃₋₇ monocyclic carbocycle or 3-7 membered monocyclic heterocycle, wherein the carbocycle or heterocycle consists of: carbon atoms and 0-2 heteroatoms selected from N, O, and S(O)_p, the carbocycle or
 30 heterocycle further comprises 0-2 double bonds and 0-2 carbonyl groups, and the carbocycle or heterocycle is substituted with 0-2 R⁴;

alternatively, Y is CY^1Y^2 , and Y^1 and Y^2 are independently C_{1-3} alkyl substituted with 0-1 R^4 ;

5 Z is selected from a bond, CH_2 , CH_2CH_2 , CH_2O , OCH_2 , $C(O)$, NH , CH_2NH , $NHCH_2$, $CH_2C(O)$, $C(O)CH_2$, $C(O)NH$, $NHC(O)$, $NHC(O)NH$, $NHC(O)CH_2C(O)NH$, $NHC(O)C(O)NH$, $C(O)NHS(O)_2$, $S(O)_2$, $CH_2S(O)_2$, $S(O)_2(CH_2)$, SO_2NH , and $NHSO_2$, provided that Z does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;

10 Z^2 is selected from H, C_{1-4} alkyl, phenyl, benzyl, $C(O)R^{3b}$, $S(O)R^{3f}$, and $S(O)_2R^{3f}$;

R^{1a} , at each occurrence, is selected from H, $-(CH_2)_r-R^{1b}$,
15 $-(CH(CH_3))_r-R^{1b}$, $-(C(CH_3)_2)_r-R^{1b}$, $-O-(CR^3R^{3a})_r-R^{1b}$, $-NR^2-(CR^3R^{3a})_r-R^{1b}$, and $-S-(CR^3R^{3a})_r-R^{1b}$, provided that R^{1a} forms other than an N-halo, N-S, O-O, or N-CN bond;

alternatively, when two R^{1a} groups are attached to adjacent
20 atoms or to the same carbon atom, together with the atoms to which they are attached they form a 5-7 membered ring consisting of: carbon atoms and 0-2 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b}
25 and 0-3 ring double bonds;

R^{1b} is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, F, Cl, Br, I, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$,
30 $NR^2C(O)NHR^2$, $NR^2C(O)_2R^{2a}$, $OC(O)NR^2R^{2a}$, $C(O)NR^2R^{2a}$, $C(O)NR^2(CH_2)_rOR^2$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle consisting of carbon atoms and from 1-4 heteroatoms

selected from the group consisting of N, O, and S(O)_p and substituted with 0-2 R^{4b}, provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;

5 R², at each occurrence, is selected from H, CF₃, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₅₋₆ carbocycle substituted with 0-2 R^{4b}, a C₅₋₆ carbocycle-CH₂- substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon
10 atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2a}, at each occurrence, is selected from H, CF₃, CH₃,
15 CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, benzyl, C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group
20 consisting of N, O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated
25 ring substituted with 0-2 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{2b}, at each occurrence, is selected from CF₃, C₁₋₄ alkoxy,
30 C₁₋₆ alkyl substituted with 0-3 R^{4b}, benzyl, C₃₋₆ carbocycle substituted with 0-2 R^{4b}, and 4-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

- R^{2c} , at each occurrence, is selected from CF_3 , OH, C_{1-4} alkoxy, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, benzyl, C_{5-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered heterocycle substituted with 0-2 R^{4b} and consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 10 R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the
- 15 group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p-S(O)_p$, S-O, O-N, O-S, or O-O moiety;
- 20 alternatively, when two R^{2d} 's are attached to the same nitrogen atom, then R^{2d} and R^{2d} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6 membered saturated, partially saturated or unsaturated ring substituted with 0-2 R^{4b} and
- 25 consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , $-(CR^3R^{3a})_r-C_{3-6}$ carbocycle
- 30 substituted with 0-2 R^{4c} , and $-(CR^3R^{3a})_r-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a C(O)-halo or C(O)- $S(O)_p$ moiety;

R^3 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

5 R^{3a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

alternatively, R^3 and R^{3a} , together with the nitrogen atom to which they are attached, combine to form a 5 or 6
10 membered saturated, partially unsaturated, or unsaturated ring consisting of: carbon atoms and the nitrogen atom to which R^3 and R^{3a} are attached;

R^{3c} , at each occurrence, is selected from CH_3 , CH_2CH_3 ,
15 $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, and phenyl;

R^{3d} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, CH_2 -phenyl, CH_2CH_2 -phenyl, and $C(=O)R^{3c}$;

20 R^{3g} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, cyclopropyl-methyl, benzyl, and phenyl;

25 alternatively, when R^3 and R^{3g} are attached to the same carbon atom, they combine with the attached carbon atom to form a cyclopropyl group;

R^4 , at each occurrence, is selected from H, =O, OR^2 , CH_2OR^2 ,
30 $(CH_2)_2OR^2$, F, Cl, Br, I, C_{1-4} alkyl, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $(CH_2)_2NR^2R^{2a}$, $C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $S(O)_pR^{5a}$, CF_3 , CF_2CF_3 , 5-6 membered carbocycle substituted with 0-1 R^5 , and a 5-6 membered heterocycle substituted with 0-1 R^5 and consisting of:

carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

5 R^{4b}, at each occurrence, is selected from H, =O, OR³,
CH₂OR³, F, Cl, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂,
CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃, C(CH₃)₃, -CN,
NO₂, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, CH₂-C(O)R³, C(O)OR^{3c},
CH₂C(O)OR^{3c}, NR³C(O)R^{3a}, CH₂NR³C(O)R^{3a}, C(O)NR³R^{3a},
CH₂C(O)NR³R^{3a}, NR³C(O)NR³R^{3a}, CH₂NR³C(O)NR³R^{3a},
10 C(=NR³)NR³R^{3a}, CH₂C(=NR³)NR³R^{3a}, NR³C(=NR³)NR³R^{3a},
CH₂NR³C(=NR³)NR³R^{3a}, SO₂NR³R^{3a}, CH₂SO₂NR³R^{3a},
NR³SO₂NR³R^{3a}, CH₂NR³SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl,
CH₂NR³SO₂-C₁₋₄ alkyl, NR³SO₂CF₃, CH₂NR³SO₂CF₃,
NR³SO₂-phenyl, CH₂NR³SO₂-phenyl, S(O)_pCF₃, CH₂S(O)_pCF₃,
15 S(O)_p-C₁₋₄ alkyl, CH₂S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl,
CH₂S(O)_p-phenyl, CF₃, and CH₂-CF₃;

R^{4c}, at each occurrence, is selected from =O, (CR³R^{3a})_rOR²,
(CR³R^{3a})_rF, (CR³R^{3a})_rBr, (CR³R^{3a})_rCl, (CR³R^{3a})_rCF₃, C₁₋₄
20 alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, (CR³R^{3a})_rCN,
(CR³R^{3a})_rNO₂, (CR³R^{3a})_rNR²R^{2a}, (CR³R^{3a})_rN(→O)R²R^{2a},
(CR³R^{3a})_rC(O)R^{2c}, (CR³R^{3a})_rNR²C(O)R^{2b},
(CR³R^{3a})_rC(O)NR²R^{2a}, (CR³R^{3a})_rNR²C(O)NR²R^{2a},
(CR³R^{3a})_rSO₂NR²R^{2a}, (CR³R^{3a})_rNR²SO₂NR²R^{2a},
25 (CR³R^{3a})_rNR²SO₂R^{5a}, (CR³R^{3a})_rS(O)_pR^{5a}, (CF₂)_rCF₃,
(CR³R^{3a})_rC₃₋₁₀ carbocycle substituted with 0-2 R^{4b}, and
(CR³R^{3a})_r5-10 membered heterocycle substituted with 0-2
R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
30 O, and S(O)_p;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$,
 $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$,
5 $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $NR^3C(O)NR^3R^{3a}$,
 $CH(=NOR^{3d})$, $C(=NR^3)NR^3R^{3a}$, $NR^3C(=NR^3)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 $NR^3SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -
phenyl, $S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 ,
10 phenyl substituted with 0-2 R^6 , naphthyl substituted
with 0-2 R^6 , and benzyl substituted with 0-2 R^6 ;

R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$,
 $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, -CN, NO_2 , NR^2R^{2a} ,
15 $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$,
 $NR^2C(O)NR^2R^{2a}$, $C(=NH)NH_2$, $NHC(=NH)NH_2$, $SO_2NR^2R^{2a}$,
 $NR^2SO_2NR^2R^{2a}$, and $NR^2SO_2C_{1-4}$ alkyl; and,

r , at each occurrence, is selected from 0, 1, 2, and 3.
20

10. A compound according to Claim 9, wherein:

25 ring M, including M_1 , M_2 , and, if present, M_3 , is selected
from phenyl, pyrrole, furan, thiophene, pyrazole,
imidazole, isoxazole, oxazole, isothiazole, thiazole,
1,2,3-triazole, 1,2,4-triazole, 1,3,4-triazole, 1,2,3-
oxadiazole, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,3-
30 thiadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole,
1,2,3,4-tetrazole, 1,2,3,5-tetrazole, pyran,
thiopyran, thiopyran-1,1-dioxide, pyridine,
pyrimidine, pyridazine, pyrazine, 1,2,3-triazine,
1,2,4-triazine, 1,2,3,4-tetrazine, dihydro-pyrrole,

5 dihydro-furan, dihydro-thiophene, dihydro-pyrazole,
dihydro-imidazole, dihydro-isoxazole, dihydro-oxazole,
dihydro-isothiazole, dihydro-thiazole, dihydro-1,2,3-
triazole, dihydro-1,2,4-triazole, dihydro-1,3,4-
triazole, dihydro-1,2,3-oxadiazole, dihydro-1,2,4-
oxadiazole, dihydro-1,3,4-oxadiazole, dihydro-1,2,3-
thiadiazole, dihydro-1,2,4-thiadiazole, dihydro-1,3,4-
thiadiazole, dihydro-1,2,3,4-tetrazole, dihydro-
1,2,3,5-tetrazole, dihydro-pyran, dihydro-thiopyran,
10 dihydro-thiopyran-1,1-dioxide, dihydro-pyridine,
dihydro-pyrimidine, dihydro-pyridazine, dihydro-
pyrazine, dihydro-1,2,3-triazine, dihydro-1,2,4-
triazine, dihydro-1,2,3,4-tetrazine, cyclopropane,
cyclobutane, cyclopentene, cyclopentane, cyclohexene,
15 cyclohexane, cycloheptane, tetrahydro-pyrrole,
tetrahydro-furan, tetrahydro-thiophene, tetrahydro-
thiophene-1,1-dioxide, tetrahydro-pyrazole,
tetrahydro-imidazole, tetrahydro-isoxazole,
tetrahydro-oxazole, tetrahydro-isothiazole,
20 tetrahydro-thiazole, tetrahydro-1,2,3-triazole,
tetrahydro-1,2,4-triazole, tetrahydro-1,3,4-triazole,
tetrahydro-1,2,3-oxadiazole, tetrahydro-1,2,4-
oxadiazole, tetrahydro-1,3,4-oxadiazole, tetrahydro-
1,2,3-thiadiazole, tetrahydro-1,2,4-thiadiazole,
25 tetrahydro-1,3,4-thiadiazole, tetrahydro-1,2,3,4-
tetrazole, tetrahydro-1,2,3,5-tetrazole, tetrahydro-
pyran, tetrahydro-thiopyran, tetrahydro-thiopyran-1,1-
dioxide, tetrahydro-pyridine, tetrahydro-pyrimidine,
tetrahydro-pyridazine, tetrahydro-pyrazine,
30 tetrahydro-1,2,3-triazine, tetrahydro-1,2,4-triazine,
tetrahydro-1,2,3,4-tetrazine, piperidine, indan,
isothiazolidine 1,1-dioxide, [1,2]thiazinane 1,1-
dioxide, 1,2,3,4-tetrahydro-naphthalene, 7,8-dimethyl-
1-oxa-spiro[4.4]nonane, 6,7-dihydro-5H-[1]pyrindine,
35 6,7-dihydro-5H-[2]pyrindine, 5,6,7,8-tetrahydro-
quinoline, 5,6,7,8-tetrahydro-isoquinoline, 5,6,7,8-

tetrahydro-quinoxaline, 6,7-dihydro-5H-cyclopentapyrazine, 4,5,6,7-tetrahydro-1H-benzoimidazole, 4,5,6,7-tetrahydro-benzothiazole, 4,5,6,7-tetrahydro-benzooxazole, 4,5,6,7-tetrahydro-5
benzo[c]isothiazole, 4,5,6,7-tetrahydro-benzo[c]isoxazole, 4,5,6,7-tetrahydro-2H-indazole, 4,5,6,7-tetrahydro-2H-isoindole, 4,5,6,7-tetrahydro-1H-indole, 5,6,7,8-tetrahydro-tetrazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-imidazo[1,2-a]pyridine, 10
4,5,6,7-tetrahydro-pyrazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-[1,2,4]triazolo[1,5-a]pyridine, 6,7-dihydro-5H-pyrrolo[1,2-c]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-a]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-b][1,2,4]triazole, 6,7-dihydro-5H-pyrrolotetrazole, 15
5,6-dihydro-4H-pyrrolo[1,2-b]pyrazole, 5,6-dihydro-4H-cyclopenta[d]isoxazole, 5,6-dihydro-4H-cyclopentaoxazole, 5,6-dihydro-4H-cyclopenta[c]isoxazole, 5,6-dihydro-4H-cyclopenta[d]isothiazole, 5,6-dihydro-4H-cyclopentathiazole, 5,6-dihydro-4H-cyclopenta[c]isothiazole, 1,4,5,6-tetrahydro-cyclopentapyrazole, 1,4,5,6-tetrahydro-cyclopentaimidazole, 2,4,5,6-tetrahydro-cyclopentapyrazole, 5,6-dihydro-4H-20
cyclopenta[1,2,5]thiadiazole, 5,6-dihydro-4H-cyclopenta[1,2,5]oxadiazole, 5,6-dihydro-4H-cyclopenta[c]furan, 2,4,5,6-tetrahydro-cyclopenta[c]pyrrole, 5,6-dihydro-4H-cyclopenta[b]furan, 5,6-dihydro-4H-30
cyclopenta[c]thiophene, 5,6-dihydro-4H-cyclopenta[b]furan, 5,6-dihydro-4H-cyclopenta[b]thiophene, 1,4,5,6-tetrahydro-cyclopenta[b]pyrrole, 2,3-dihydro-1H-indolizin-5-one, 6,7,8,9-tetrahydro-quinolizin-4-one, 1-oxa-35
spiro[4.4]nonane, 1-aza-spiro[4.4]nonane, 2-oxa-spiro[4.4]nonane, 2-aza-spiro[4.4]nonane, 1-aza-

spiro[4.5]decane, 1-oxa-spiro[4.5]decane, 2-oxa-
 spiro[4.5]decane, 2-aza-spiro[4.5]decane, 1-thia-
 spiro[4.4]nonane, 1-thia-spiro[4.5]decane, 2-thia-
 spiro[4.4]nonane, 2-thia-spiro[4.5]decane, 7-oxa-
 5 bicyclo[2.2.1]heptane, 2-oxa-bicyclo[2.2.1]heptane, 7-
 thia-bicyclo[2.2.1]heptane, 2-thia-
 bicyclo[2.2.1]heptane, 2-aza-bicyclo[2.2.1]heptane, 7-
 aza-bicyclo[2.2.1]heptane, 4,5,6,7-tetrahydro-
 benzo[d]isoxazole, 4,5,6,7-tetrahydro-benzooxazole,
 10 4,5,6,7-tetrahydro-benzo[d]isothiazole, 4,5,6,7-
 tetrahydro-benzothiazole, 4,5,6,7-tetrahydro-1H-
 indazole, 4,5,6,7-tetrahydro-benzo[c]thiophene,
 4,5,6,7-tetrahydro-benzo[b]thiophene, 4,5,6,7-
 tetrahydro-isobenzofuran, 4,5,6,7-tetrahydro-
 15 benzofuran, 5,6,7,8-tetrahydro-quinoxaline, 6,7-
 dihydro-5H-cyclopentapyrazine, 5,6,7,8-tetrahydro-
 imidazo[1,5-a]pyridine, 5,6,7,8-tetrahydro-
 imidazo[1,2-a]pyridine, 5,6,7,8-tetrahydro-
 [1,2,4]triazolo[1,5-a]pyridine, 5,6,7,8-tetrahydro-
 20 tetrazolo[1,5-a]pyridine, 4,5,6,7-tetrahydro-
 pyrazolo[1,5-a]pyridine, 6,7-dihydro-5H-pyrrolo[1,2-
 a]imidazole, 6,7-dihydro-5H-pyrrolo[1,2-
 b][1,2,4]triazole, 5,6-dihydro-4H-pyrrolo[1,2-
 b]pyrazole, and 6,7-dihydro-5H-pyrrolotetrazole;

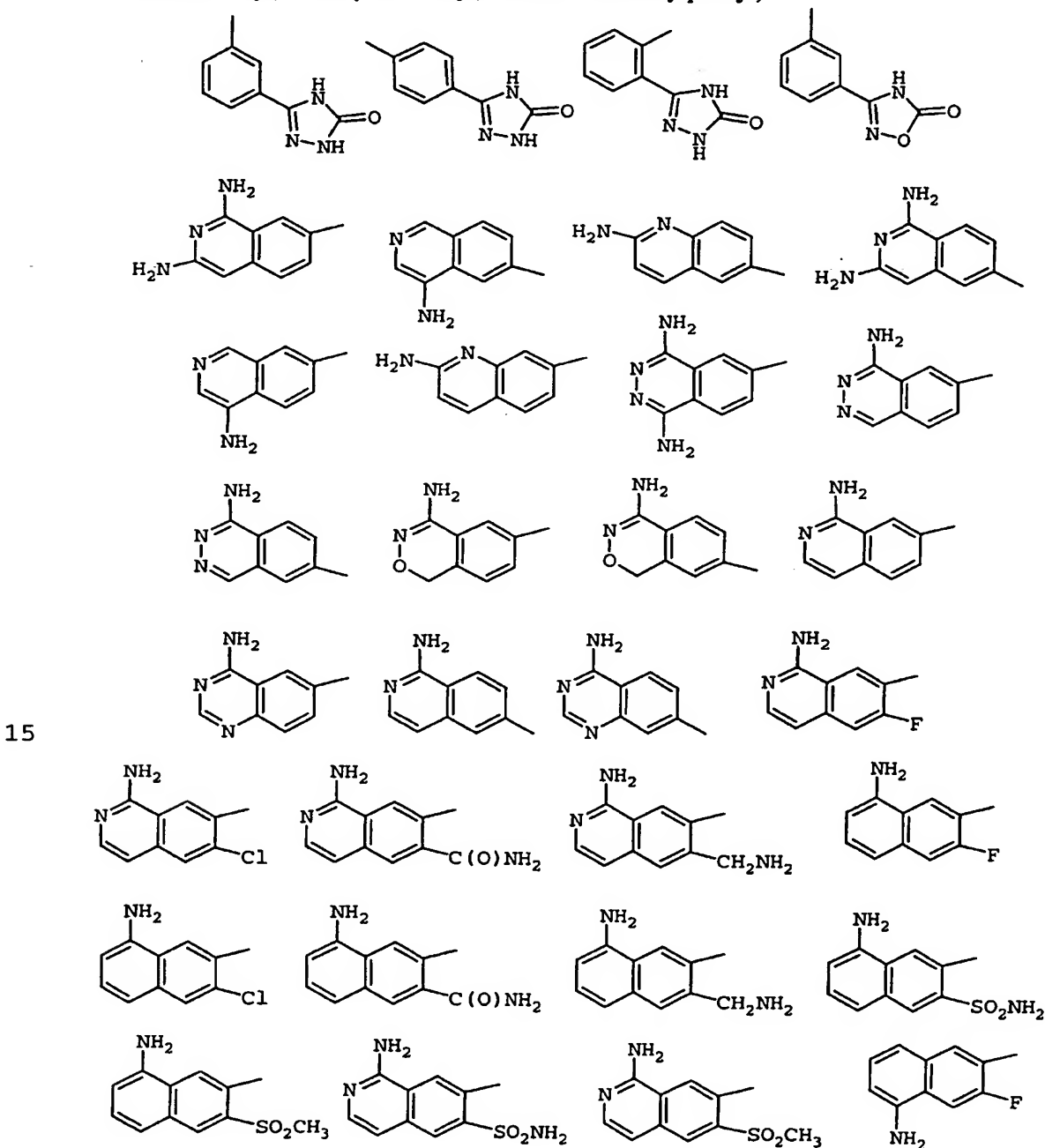
25

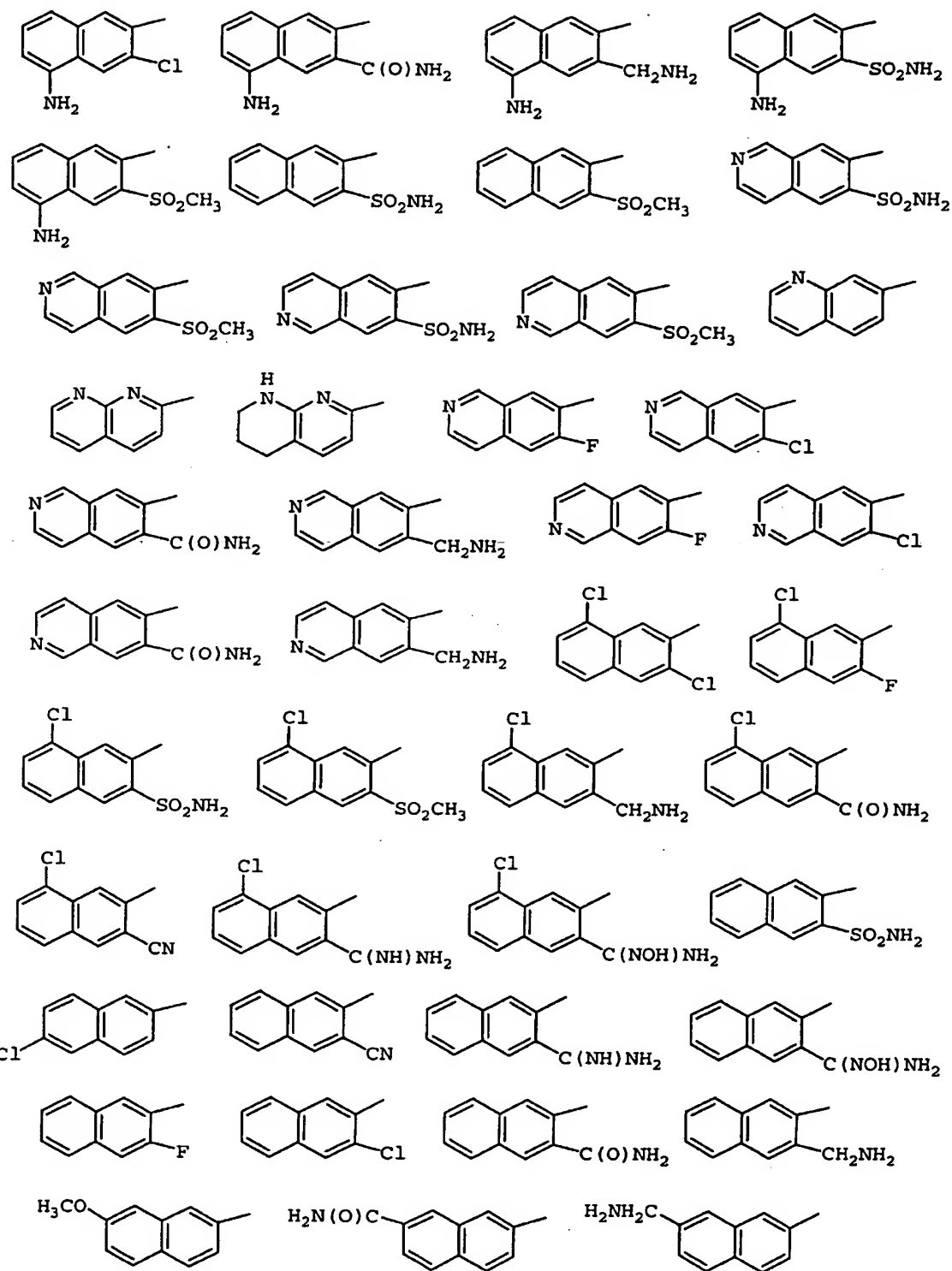
ring M is substituted with 0-3 R^{1a} and 0-1 carbonyl group;

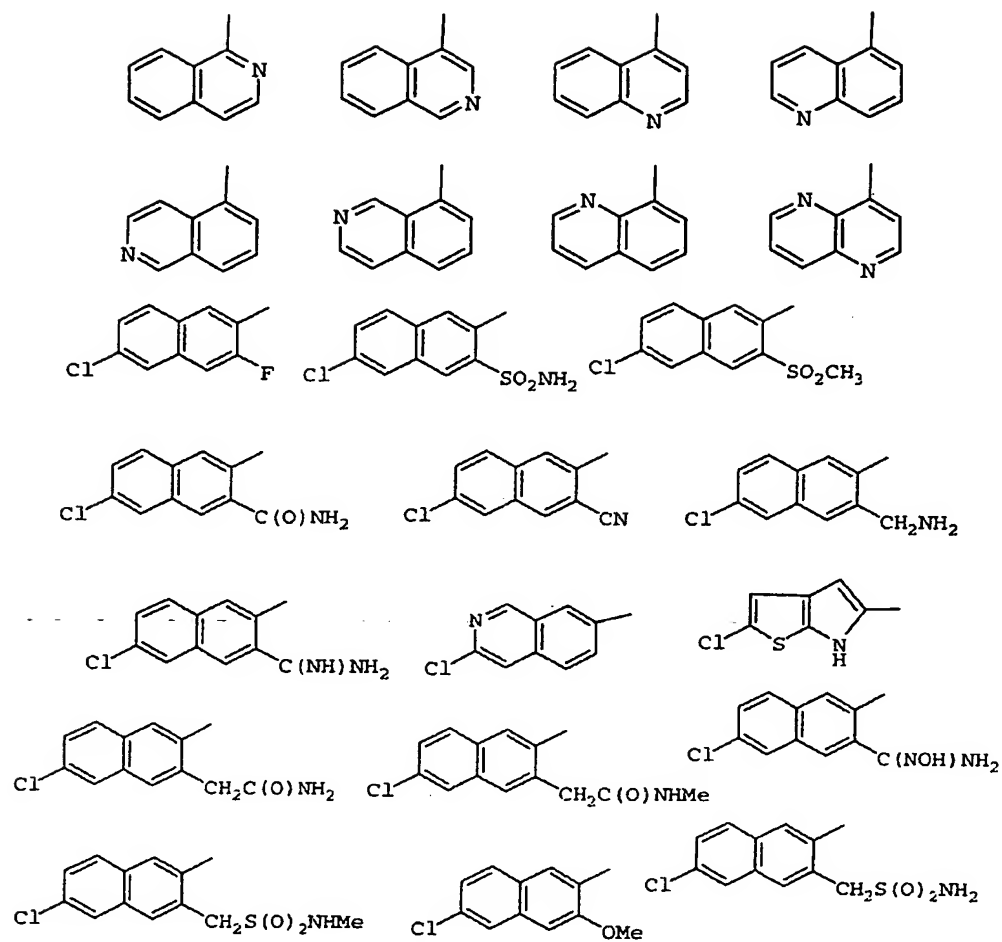
G is selected from the group:

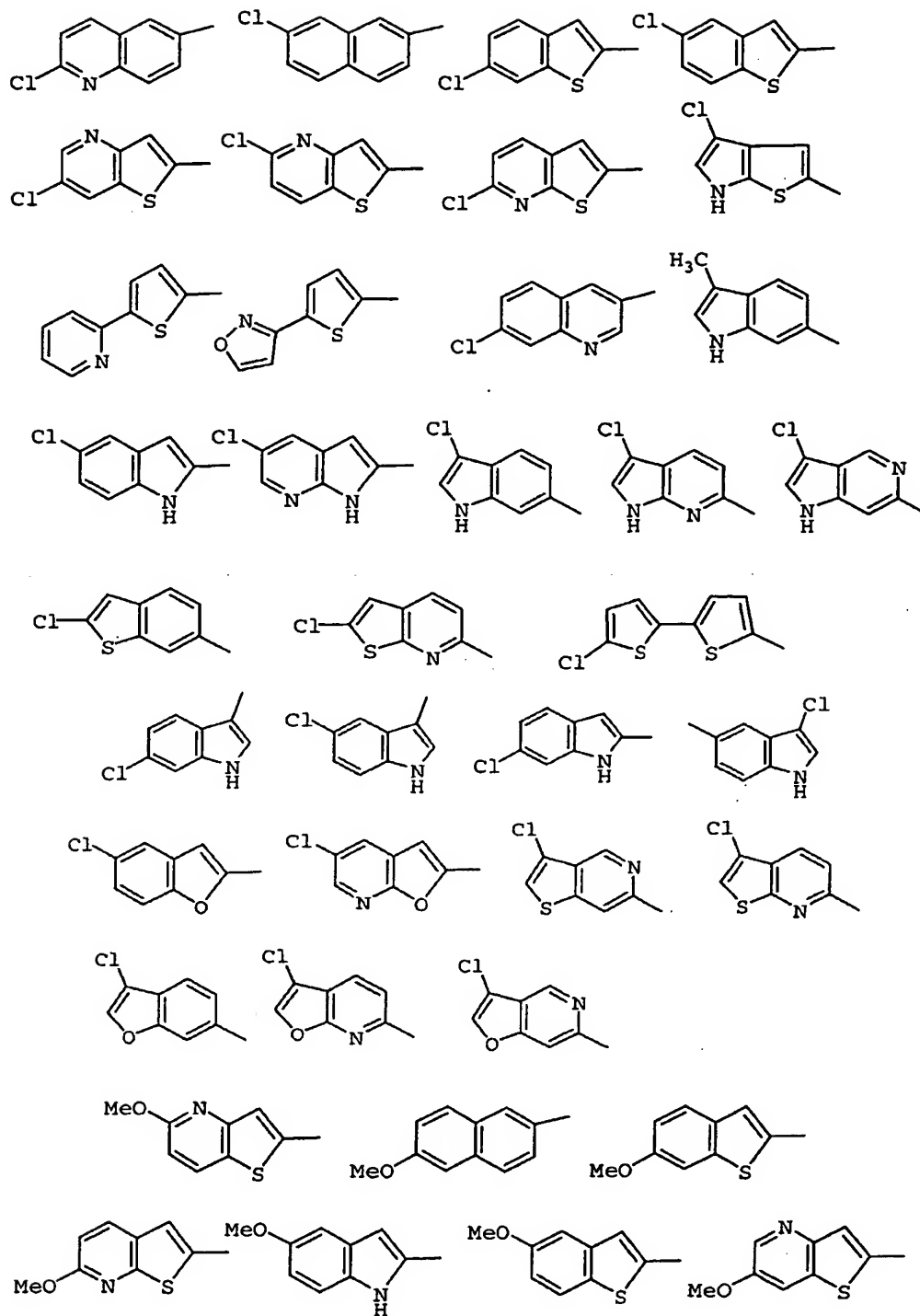
phenyl; 4-ethyl-phenyl; 2,5-bis-aminomethyl-phenyl; 2-amido-4-methoxy-phenyl;
 30 2-amido-5-chloro-phenyl; 2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
 2-aminomethyl-3-methoxy-phenyl; 2-aminomethyl-4-fluoro-phenyl;
 2-aminomethyl-4-methoxy-phenyl; 2-aminomethyl-5-fluoro-phenyl;
 2-aminomethyl-5-methoxy-phenyl; 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl; 2-aminosulfonyl-phenyl;
 35 2-hydroxy-4-methoxy-phenyl; 2-methylsulfonyl-phenyl; 3-(N,N-dimethylamino)-4-chloro-phenyl;
 3-(N,N-dimethylamino)-phenyl; 3-(N-hydroxy-amidino)-phenyl; 3-(N-methoxy-amidino)-phenyl;
 3-(N-methylamino)-4-chloro-phenyl; 3-(N-methylamino)-phenyl; 3-amidino-phenyl;
 3-amido-6-hydroxy-phenyl; 3-amido-phenyl; 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
 3-amino-phenyl; 3-chloro-4-fluoro-phenyl; 3-chloro-phenyl; 3-hydroxy-4-methoxy-phenyl; 3,5-
 40 dichloro-thien-2-yl; 4-(N,N-dimethylamino)-5-chloro-thien-2-yl;
 4-(N-methylamino)-5-chloro-thien-2-yl; 4-amino-5-chloro-thien-2-yl; 4-amino-pyrid-2-yl;
 4-chloro-3-fluoro-phenyl; 4-chloro-phenyl; 4-chloro-pyrid-2-yl; 4-methoxy-2-methylsulfonyl-phenyl;

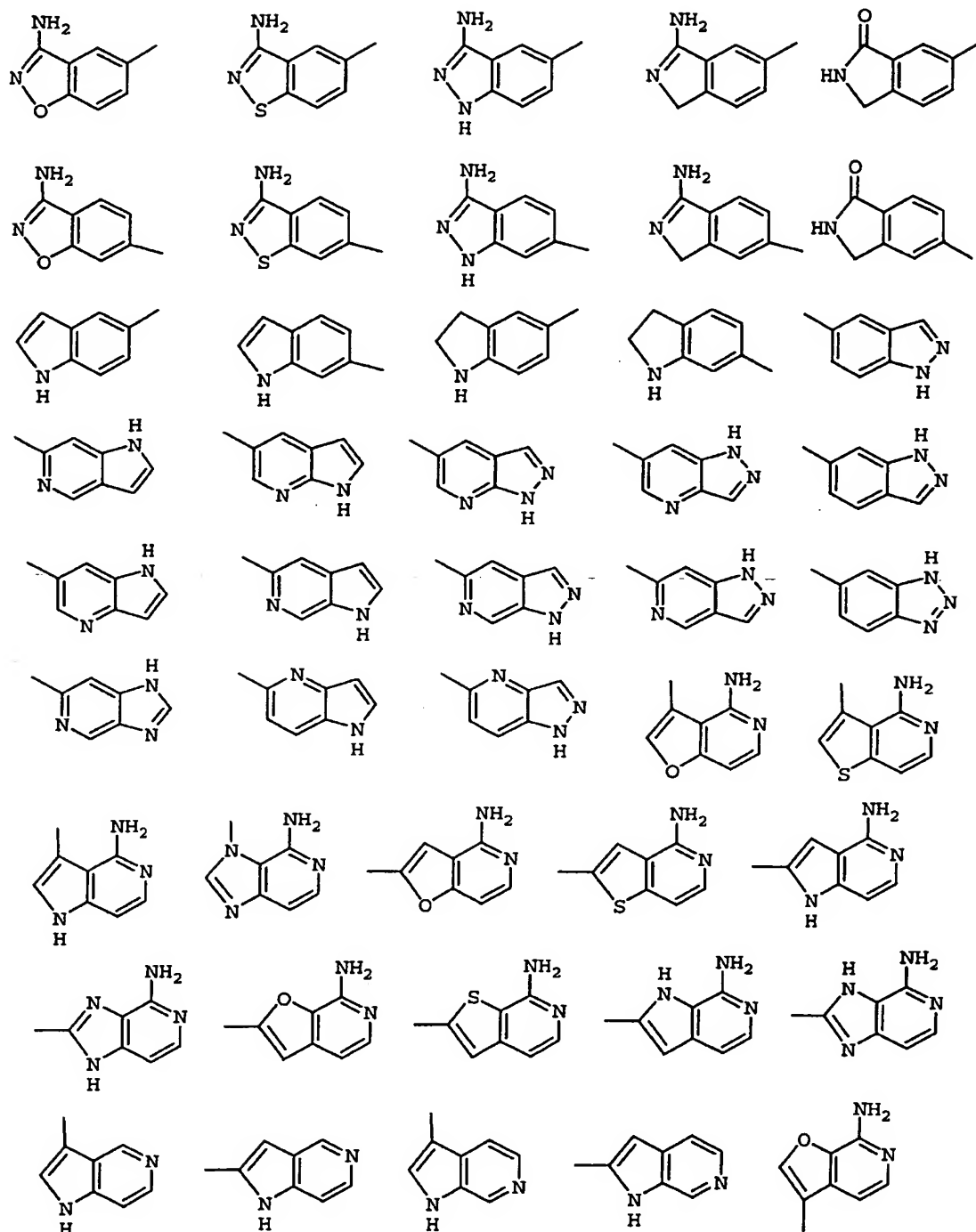
4-methoxy-phenyl; 2-methoxy-pyrid-5-yl; 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;
5-(N-methylamino)-4-chloro-thien-2-yl; 5-amino-4-chloro-thien-2-yl;
5-chloro-2-aminosulfonyl-phenyl; 5-chloro-2-methylsulfonyl-phenyl; 5-chloro-pyrid-2-yl;
5-chloro-thien-2-yl; 5-methoxy-thien-2-yl; 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-
chloro-pyrimidin-3-yl; 6-chloro-pyridazin-3-yl; 2-aminomethyl-4-chloro-phenyl;
2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl; 4-chloro-2-methylsulfonyl-phenyl;
2-aminosulfonyl-4-fluoro-phenyl; 2-amido-4-fluoro-phenyl; 4-fluoro-2-methylsulfonyl-phenyl;
2-aminomethyl-4-bromo-phenyl; 2-aminosulfonyl-4-bromo-phenyl; 2-amido-4-bromo-phenyl;
4-bromo-2-methylsulfonyl-phenyl; 2-aminomethyl-4-methyl-phenyl;
2-aminosulfonyl-4-methyl-phenyl; 2-amido-4-methyl-phenyl; 2-methylsulfonyl-4-methyl-phenyl;
4-fluoro-pyrid-2-yl; 4-bromo-pyrid-2-yl; 4-methyl-pyrid-2-yl; 5-fluoro-thien-2-yl;
5-bromo-thien-2-yl; 5-methyl-thien-2-yl; 2-amido-4-methoxy-phenyl;

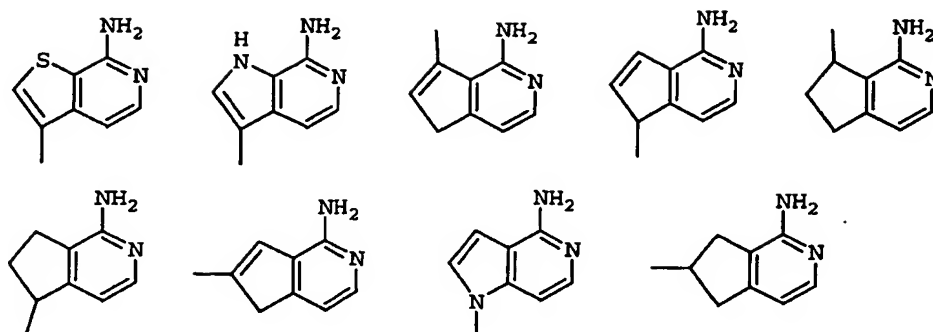












;

G_1 is absent or is selected from $(CR^3R^{3a})_{1-3}$, $CR^3=CR^3$,

$(CR^3R^{3a})_u C(O) (CR^3R^{3a})_w$, $(CR^3R^{3a})_u O (CR^3R^{3a})_w$,

5 $(CR^3R^{3a})_u NR^{3b} (CR^3R^{3a})_w$, $(CR^3R^{3a})_u C(O) NR^{3b} (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u NR^{3b} C(O) (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u NR^{3b} C(O) (CR^3R^{3a})_u C(O) NR^{3b} (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u S (CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O) (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u S(O)_2 (CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O) NR^{3b} (CR^3R^{3a})_w$,

10 $(CR^3R^{3a})_u NR^{3b} S(O)_2 (CR^3R^{3a})_w$, $(CR^3R^{3a})_u S(O)_2 NR^{3b} (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u C(O) NR^{3b} S(O)_2 (CR^3R^{3a})_w$,

$(CR^3R^{3a})_u NR^{3b} C(S) (CR^3R^{3a})_u C(O) NR^{3b} (CR^3R^{3a})_w$, and

$(CR^3R^{3a})_u NR^{3b} C(O) (CR^3R^{3a})_u C(S) NR^{3b} (CR^3R^{3a})_w$, wherein u

+ w total 0, 1, or 2, provided that G_1 does not form a

15 N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

A is selected from one of the following carbocycles and heterocycles which are substituted with 0-2 R^4 ;

20 cyclohexyl, phenyl, piperidinyl, piperazinyl, pyridyl, pyrimidyl, furanyl, morpholinyl, thienyl, pyrrolyl, pyrrolidinyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyrazolyl, imidazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl,

25 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl,

1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl,
1,3,4-triazolyl, benzofuranyl, benzothiofuranyl,
indolynyl, indolyl, benzimidazolyl, benzoxazolyl,
benzthiazolyl, indazolyl, benzisoxazolyl,
5 benzisothiazolyl, and isoindazolyl;

X is selected from $-(CR^2R^{2a})_{1-2}-$, $-C(O)-$, $-S(O)_2-$,
 $-NR^2S(O)_2-$, $-NR^2S(O)_2NR^2-$, $-NR^2C(O)-$, $-C(O)NR^2-$, NR^2 ,
10 $-NR^2CR^2R^{2a}-$, $-CR^2R^{2a}NR^2-$, O, $-OCR^2R^{2a}-$, and $-CR^2R^{2a}O-$;

10

Y is a C_{3-6} monocyclic carbocycle or 5-6 membered monocyclic
heterocycle, wherein the carbocycle or heterocycle
consists of carbon atoms and 0-2 heteroatoms selected
from N, O, and $S(O)_p$, the carbocycle or heterocycle
15 further comprises 0-1 double bonds and 0-1 carbonyl
groups, and the carbocycle or heterocycle is
substituted with 0-2 R^4 ;

alternatively, Y is CY^1Y^2 , and Y^1 and Y^2 are independently
20 C_{1-2} alkyl substituted with 0-1 R^4 ;

R^{1a} , at each occurrence, is selected from H, R^{1b} ,
 $CH(CH_3)R^{1b}$, $C(CH_3)_2R^{1b}$, CH_2R^{1b} , and $CH_2CH_2R^{1b}$, provided
that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

25

alternatively, when two R^{1a} groups are attached to adjacent
atoms or to the same carbon atom, together with the
atoms to which they are attached, they form a 5-6
membered ring consisting of: carbon atoms and 0-2
30 heteroatoms selected from the group consisting of N,
O, and $S(O)_p$, this ring being substituted with 0-2 R^{4b}
and comprising: 0-3 double bonds;

- R^{1b} is selected from H, CH_3 , CH_2CH_3 , F, Cl, Br, -CN, -CHO, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , $OC(O)R^2$, CO_2R^{2a} , $S(O)_pR^2$, $NR^2(CH_2)_rOR^2$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and
 5 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;
- 10 R^2 , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-2 R^{4b} , benzyl substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b}
 15 and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 20 R^{2a} , at each occurrence, is selected from H, CF_3 , CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 25 R^{2b} , at each occurrence, is selected from CF_3 , C_{1-4} alkoxy, C_{1-5} alkyl substituted with 0-3 R^{4b} , benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 4-6 membered substituted with 0-2 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group
 30 consisting of N, O, and $S(O)_p$;
- R^{2c} , at each occurrence, is selected from CF_3 , OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$,

CH(CH₃)₂, benzyl, phenyl substituted with 0-2 R^{4b}, and
5-6 membered aromatic heterocycle substituted with 0-2
R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
5 O, and S(O)_p;

alternatively, R² and R^{2a}, together with the nitrogen atom
to which they are attached, combine to form a 3-6
membered saturated, partially saturated or unsaturated
10 ring substituted with 0-2 R^{4b} and consisting of: 0-1
additional heteroatoms selected from the group
consisting of N, O, and S(O)_p;

R^{2d}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl
15 substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted
with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with
0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2
R^{4c} and consisting of: carbon atoms and 1-4
heteroatoms selected from the group consisting of N,
20 O, and S(O)_p, and -(CR³R^{3a})-5-6 membered heterocycle
substituted with 0-2 R^{4c} and consisting of: carbon
atoms and 1-4 heteroatoms selected from the group
consisting of N, O, and S(O)_p, provided that R^{2d} forms
other than a N-halo, N-C-halo, S(O)_p-halo, O-halo, N-
25 S, S-N, S(O)_p-S(O)_p, S-O, O-N, O-S, or O-O moiety;

R^{2e}, at each occurrence, is selected from H, R^{4c}, C₁₋₄ alkyl
substituted with 0-2 R^{4c}, C₃₋₆ carbocycle substituted
with 0-2 R^{4c}, -(CR³R^{3a})-C₃₋₆ carbocycle substituted with
30 0-2 R^{4c}, 5-6 membered heterocycle substituted with 0-2
R^{4c} consisting of: carbon atoms and 1-4 heteroatoms
selected from the group consisting of N, O, and S(O)_p,
and -(CR³R^{3a})-5-6 membered heterocycle substituted with
0-2 R^{4c} and consisting of: carbon atoms and 1-4

heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a $C(O)$ -halo or $C(O)-S(O)_p$ moiety;

5 R^4 , at each occurrence, is selected from H, $(CH_2)_2OR^2$, CH_2OR^2 , OR^2 , F, Cl, Br, I, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, $CH_2CH_2CH_2CH_3$, $CH_2CH(CH_3)_2$, $CH(CH_3)CH_2CH_3$, $C(CH_3)_3$, -CN, NO_2 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $(CH_2)_2NR^2R^{2a}$, $C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, CF_3 , and
10 CF_2CF_3 ;

R^{4a} is selected from $-(CR^3R^{3g})_r$ -5-6 membered carbocycle substituted with 0-3 R^{4c} , $-(CR^3R^{3g})_r$ -5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of:
15 carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, $(CR^3R^{3g})_rNR^{2d}R^{2d}$, $(CR^3R^{3g})_rN(\rightarrow O)R^{2d}R^{2d}$, $(CR^3R^{3g})_rOR^{2d}$, $(CR^3R^{3g})_r-NR^{2d}C(O)R^{2e}$, $(CR^3R^{3g})_r-C(O)R^{2e}$, $(CR^3R^{3g})_r-OC(O)R^{2e}$, $(CR^3R^{3g})_r-C(O)NR^{2d}R^{2d}$,
20 $(CR^3R^{3g})_r-C(O)OR^{2d}$, $(CR^3R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d}$, $(CR^3R^{3g})_r-NR^{2d}C(O)OR^{2d}$, $(CR^3R^{3g})_r-SO_2NR^{2d}R^{2d}$, $(CR^3R^{3g})_r-NR^{2d}SO_2R^{2d}$, and $(CR^3R^{3g})_r-S(O)_pR^{2d}$, provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or $S(O)H$;

25 R^{4b} , at each occurrence, is selected from H, $=O$, OR^3 , CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN, NO_2 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2-C(O)R^3$, $C(O)OR^{3c}$, $CH_2-C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $CH_2NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $CH_2-C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$, $CH_2SO_2NR^3R^{3a}$, $NR^3SO_2-C_{1-4}$ alkyl,
30 $CH_2NR^3SO_2-C_{1-4}$ alkyl, NR^3SO_2 -phenyl, $CH_2NR^3SO_2$ -phenyl, $S(O)_pCF_3$, $CH_2S(O)_pCF_3$, $S(O)_p-C_{1-4}$ alkyl, $CH_2S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, $CH_2S(O)_p$ -phenyl, and CF_3 ;

R^{4c} , at each occurrence, is selected from =O, OR^2 ,
 $(CR^3R^{3a})OR^2$, F, $(CR^3R^{3a})F$, Br, $(CR^3R^{3a})Br$, Cl,
 $(CR^3R^{3a})Cl$, CF_3 , $(CR^3R^{3a})CF_3$, C_{1-4} alkyl, C_{2-3} alkenyl,
5 C_{2-3} alkynyl, -CN, $(CR^3R^{3a})CN$, NO_2 , $(CR^3R^{3a})NO_2$, NR^2R^{2a} ,
 $(CR^3R^{3a})NR^2R^{2a}$, $N(\rightarrow O)R^2R^{2a}$, $(CR^3R^{3a})N(\rightarrow O)R^2R^{2a}$, $C(O)R^{2c}$,
 $(CR^3R^{3a})C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $(CR^3R^{3a})NR^2C(O)R^{2b}$,
 $C(O)NR^2R^{2a}$, $(CR^3R^{3a})C(O)NR^2R^{2a}$, $NR^2C(O)NR^2R^{2a}$,
 $(CR^3R^{3a})NR^2C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $(CR^3R^{3a})SO_2NR^2R^{2a}$,
10 $NR^2SO_2NR^2R^{2a}$, $(CR^3R^{3a})NR^2SO_2NR^2R^{2a}$, $NR^2SO_2R^{5a}$,
 $(CR^3R^{3a})NR^2SO_2R^{5a}$, $S(O)_pR^{5a}$, $(CR^3R^{3a})S(O)_pR^{5a}$, CF_3 ,
 CF_2CF_3 , C_{3-10} carbocycle substituted with 0-2 R^{4b} ,
 $(CR^3R^{3a})C_{3-10}$ carbocycle substituted with 0-2 R^{4b} , 5-10
membered heterocycle substituted with 0-2 R^{4b} and
15 consisting of carbon atoms and from 1-4 heteroatoms
selected from the group consisting of N, O, and $S(O)_p$,
and (CR^3R^{3a}) 5-10 membered heterocycle substituted with
0-2 R^{4b} and consisting of carbon atoms and from 1-4
heteroatoms selected from the group consisting of N,
20 O, and $S(O)_p$;

R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $CH_2CH_2CH_3$, $CH(CH_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$, $CH_2C(O)R^3$, $C(O)OR^{3c}$,
25 $CH_2C(O)OR^{3c}$, $NR^3C(O)R^{3a}$, $C(O)NR^3R^{3a}$, $SO_2NR^3R^{3a}$,
 $NR^3SO_2-C_{1-4}$ alkyl, $NR^3SO_2CF_3$, NR^3SO_2 -phenyl, $S(O)_pCF_3$,
 $S(O)_p-C_{1-4}$ alkyl, $S(O)_p$ -phenyl, CF_3 , phenyl substituted
with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
benzyl substituted with 0-2 R^6 ;

30

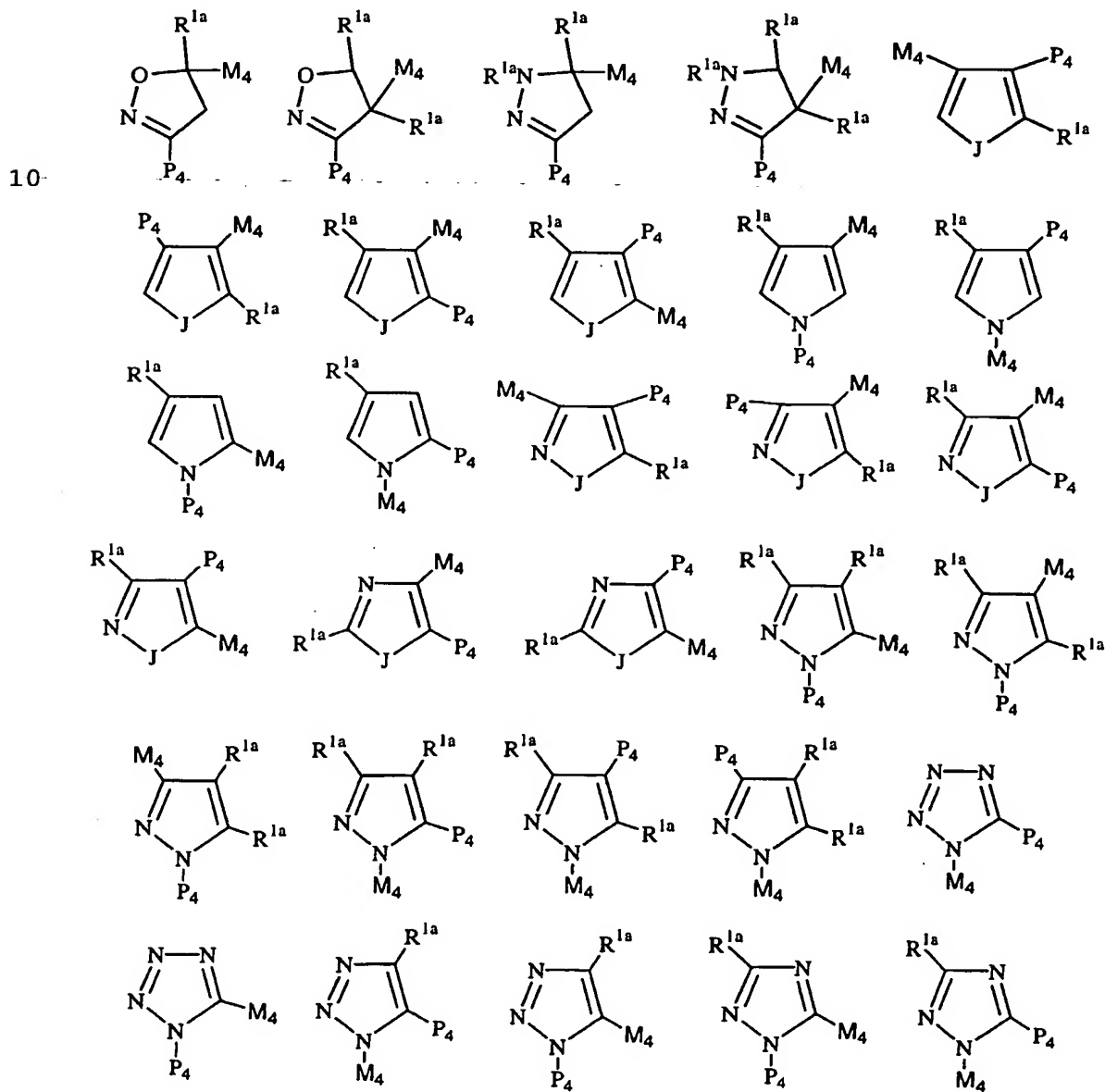
R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, -CN, NO_2 , NR^2R^{2a} ,

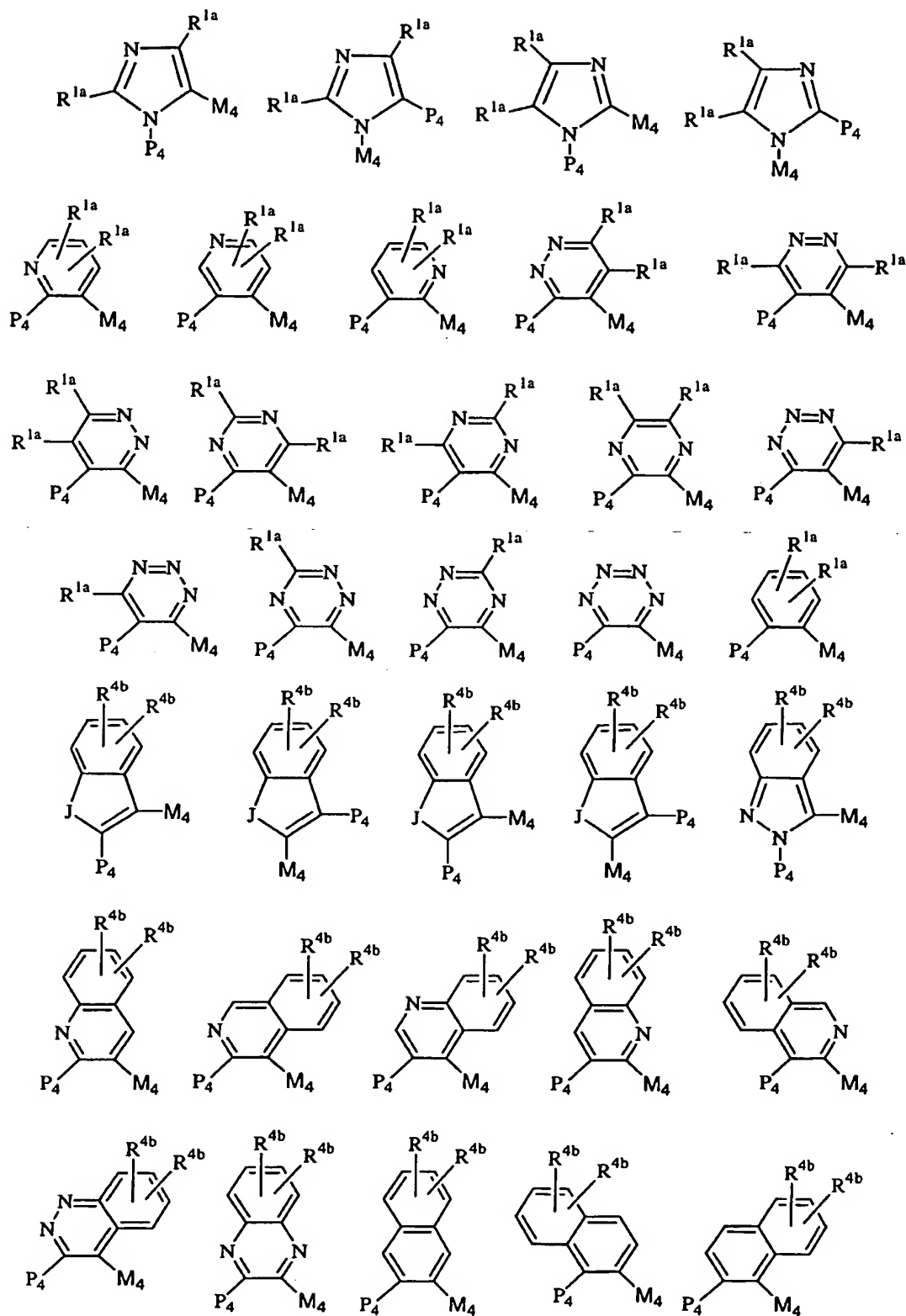
$\text{CH}_2\text{NR}^2\text{R}^{2a}$, $\text{C}(\text{O})\text{R}^{2b}$, $\text{CH}_2\text{C}(\text{O})\text{R}^{2b}$, $\text{NR}^2\text{C}(\text{O})\text{R}^{2b}$, $\text{SO}_2\text{NR}^2\text{R}^{2a}$,
and $\text{NR}^2\text{SO}_2\text{C}_{1-4}$ alkyl; and,

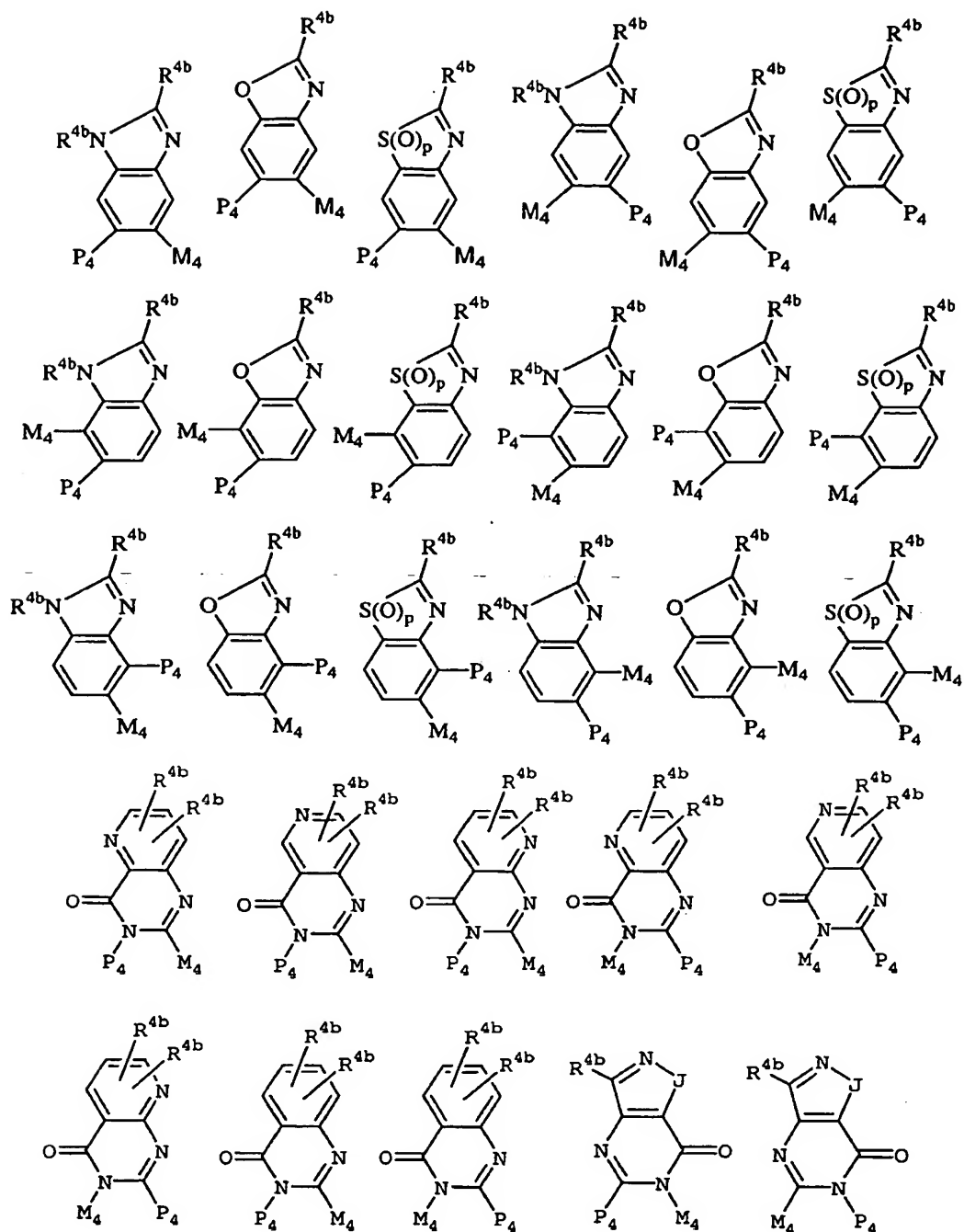
r , at each occurrence, is selected from 0, 1, and 2.

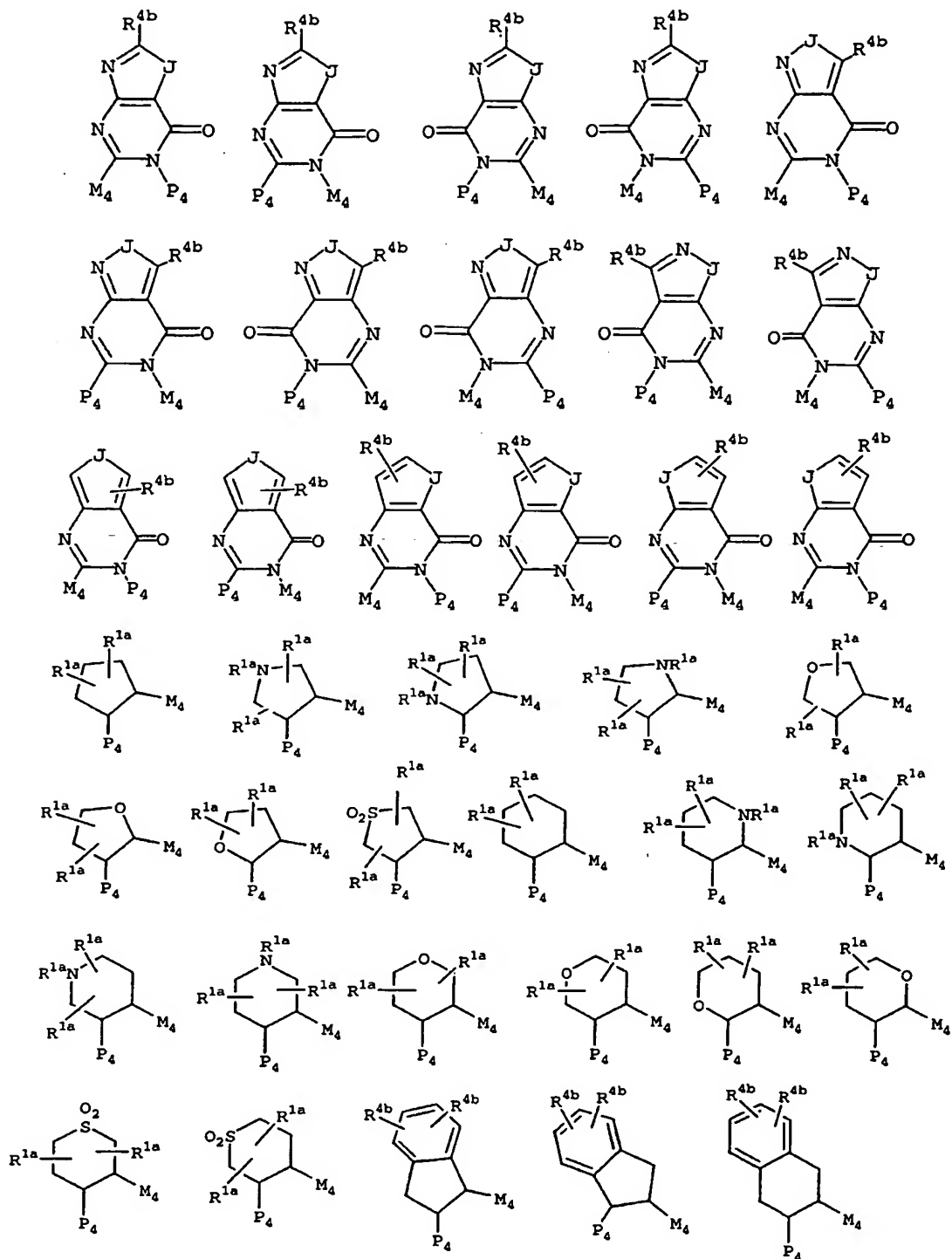
5

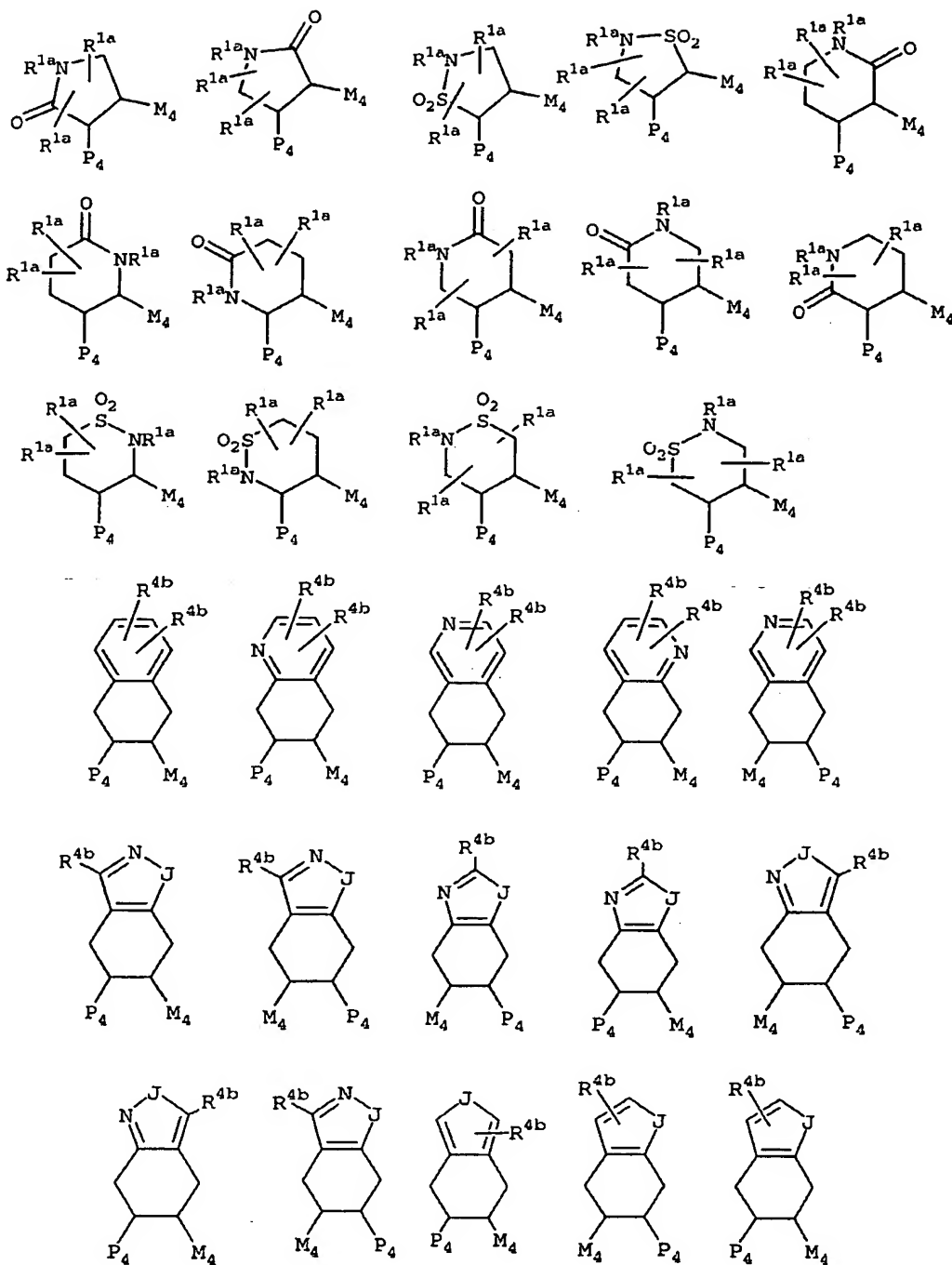
11. A compound according to Claim 10, wherein: the compound is selected from:

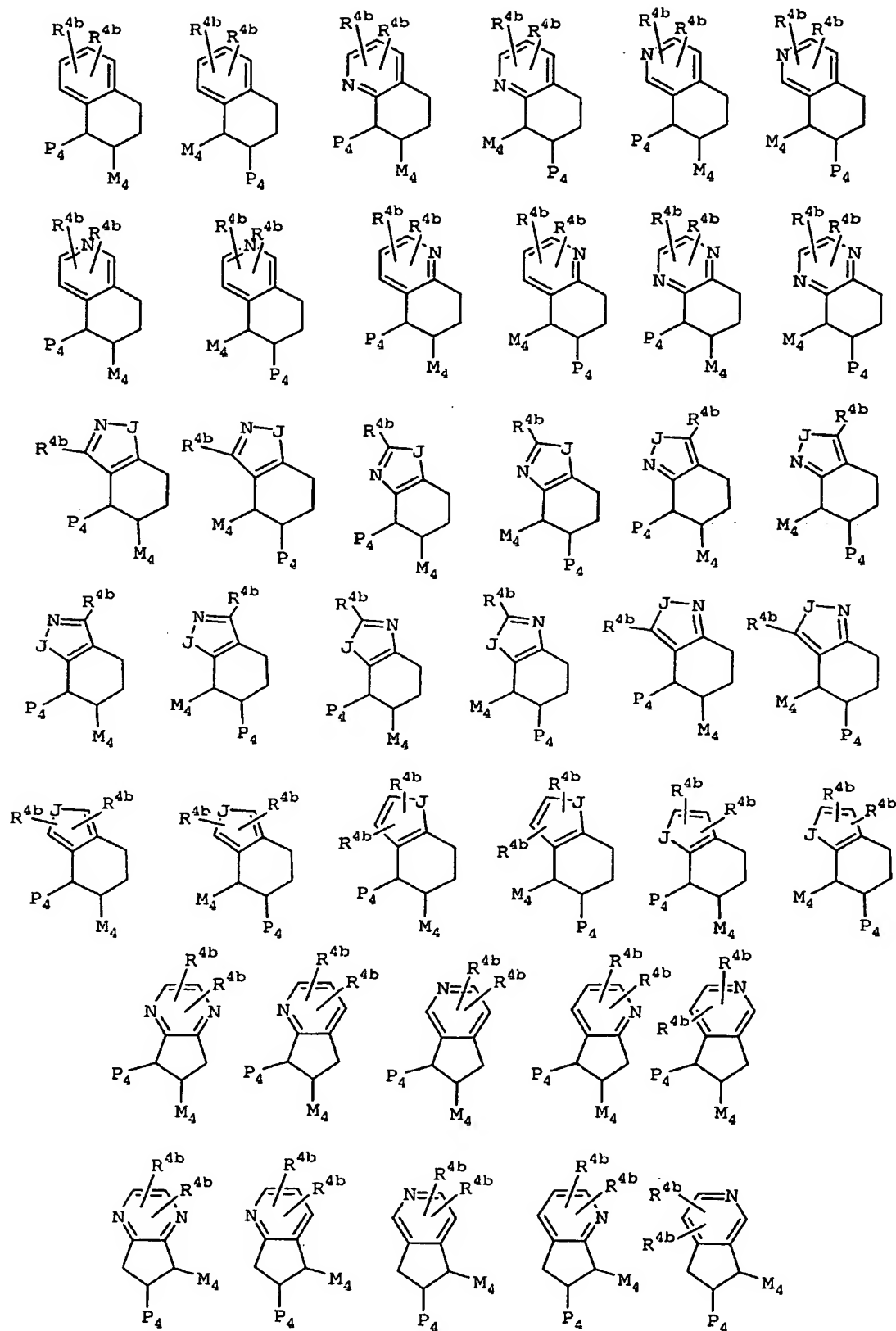


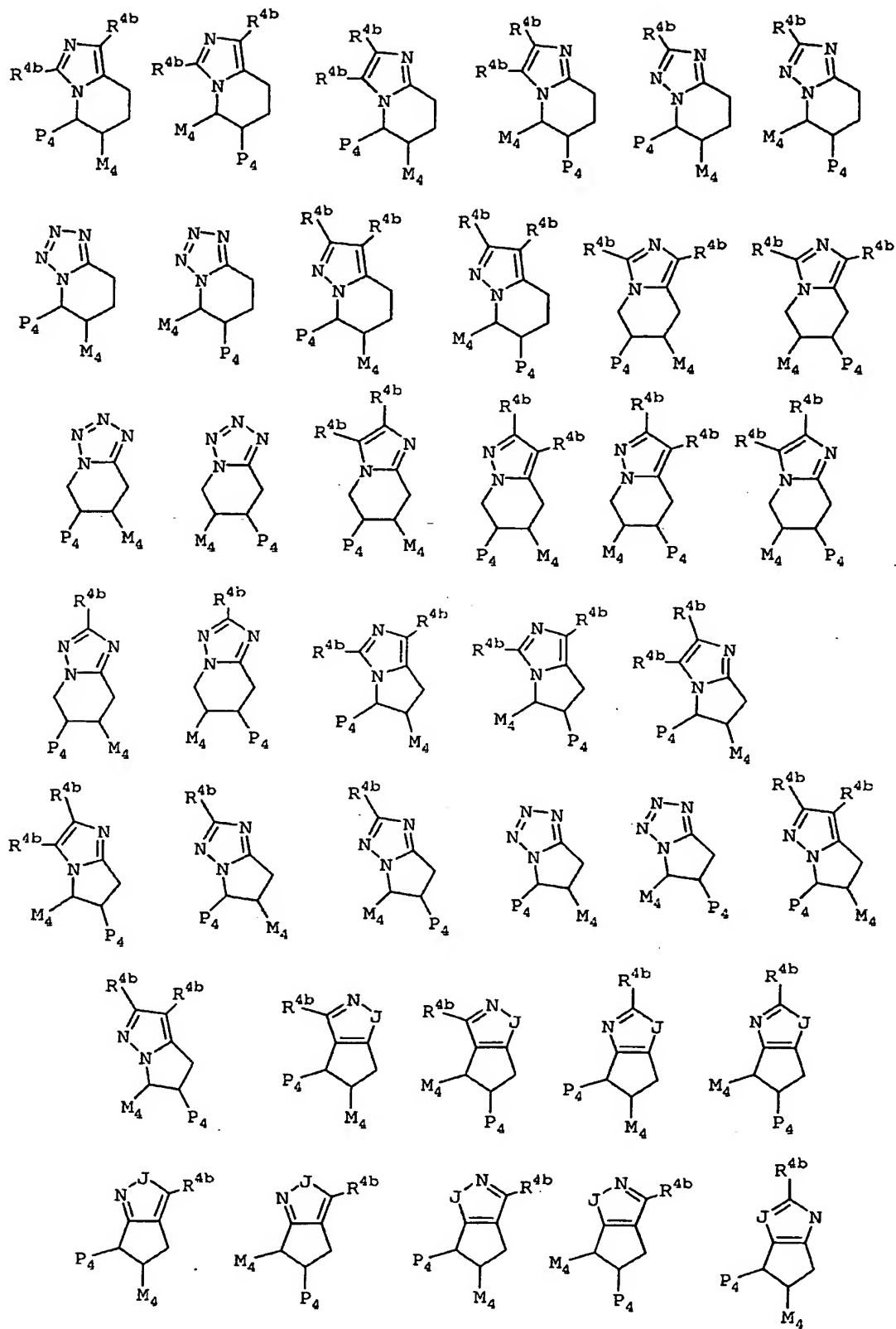


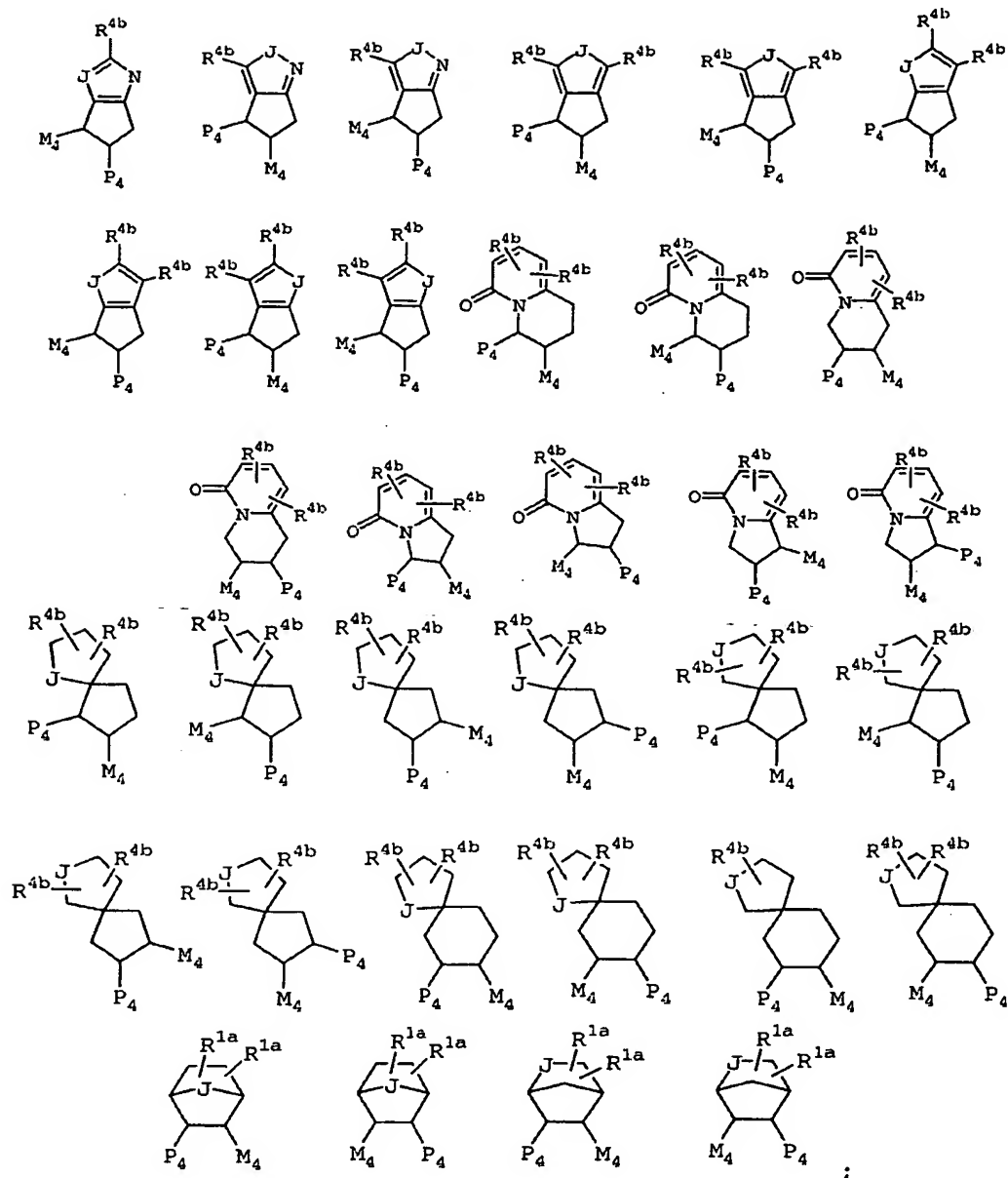










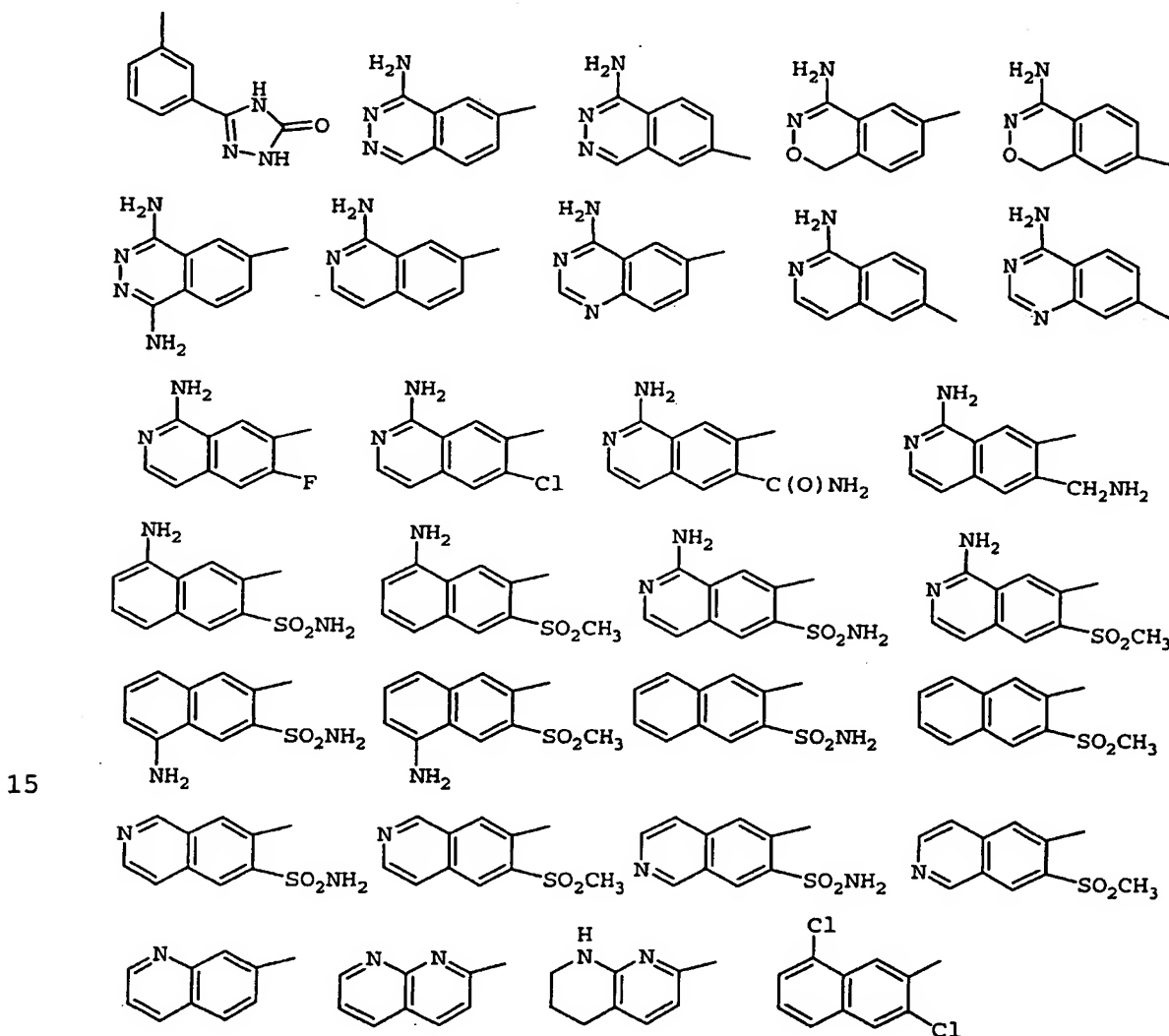


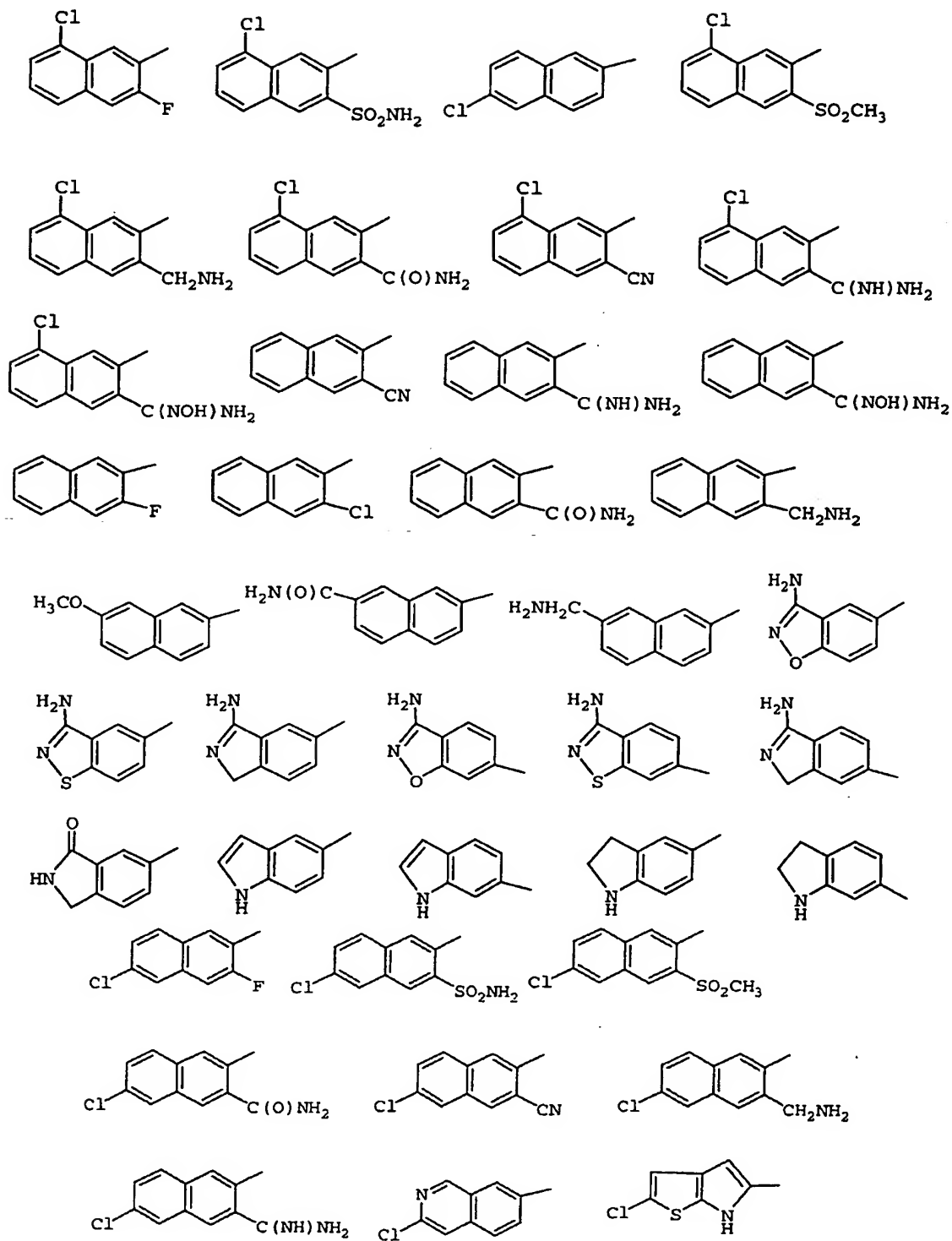
5 J is selected from O, S, NH, and NR^{1a};

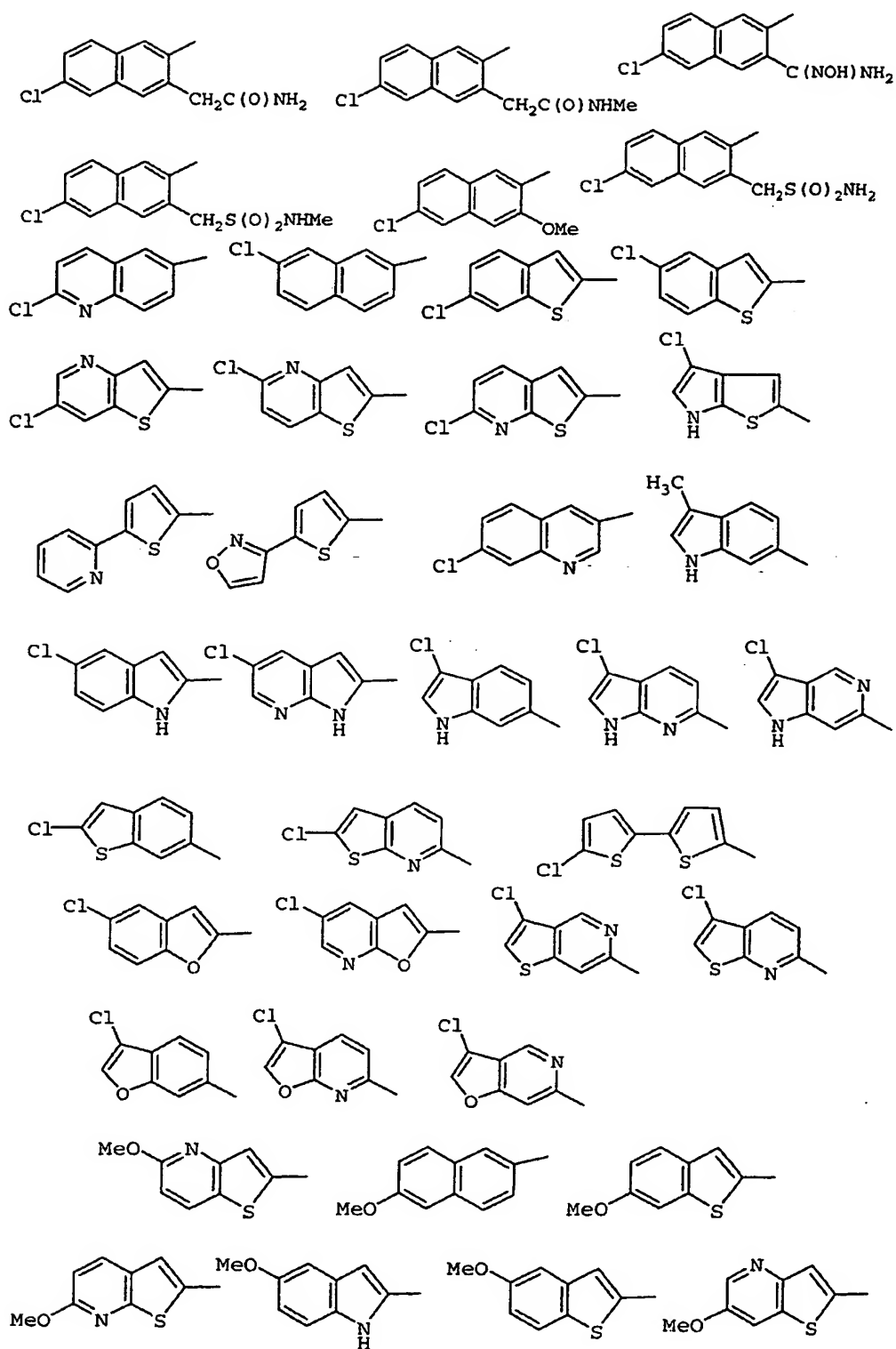
G is selected from the group:

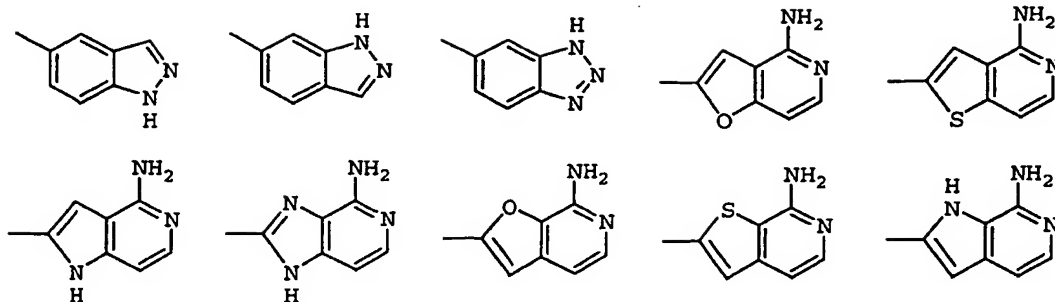
- 2-amido-4-methoxy-phenyl; 2-amido-phenyl; 2-aminomethyl-3-fluoro-phenyl;
 2-aminomethyl-4-fluoro-phenyl; 2-aminomethyl-4-methoxy-phenyl; 2-aminomethyl-5-fluoro-phenyl;
 10 2-aminomethyl-5-methoxy-phenyl; 2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl; 2-aminosulfonyl-phenyl;
 2-methylsulfonyl-phenyl; 3-(N,N-dimethylamino)-4-chloro-phenyl; 3-(N,N-dimethylamino)-phenyl;
 3-(N-methylamino)-4-chloro-phenyl; 3-(N-methylamino)-phenyl; 3-amido-phenyl;
 3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl; 3-amino-phenyl; 3-chloro-phenyl; 3,5-dichloro-
 15 thien-2-yl; 4-(N,N-dimethylamino)-5-chloro-thien-2-yl; 4-(N-methylamino)-5-chloro-thien-2-yl;
 4-amino-5-chloro-thien-2-yl; 4-chloro-phenyl; 4-methoxy-2-methylsulfonyl-phenyl;
 4-methoxy-phenyl; 5-(N,N-dimethylamino)-4-chloro-thien-2-yl;

- 5-(N-methylamino)-4-chloro-thien-2-yl; 5-amino-4-chloro-thien-2-yl; 5-chloro-pyrid-2-yl;
 5-chloro-thien-2-yl; 5-methoxy-thien-2-yl; 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-
 chloro-pyrimidin-3-yl; 6-chloro-pyridazin-3-yl; 2-aminomethyl-4-chloro-phenyl;
 2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl; 4-chloro-2-methylsulfonyl-phenyl;
 5
 2-aminosulfonyl-4-fluoro-phenyl; 2-amido-4-fluoro-phenyl; 4-fluoro-2-methylsulfonyl-phenyl;
 2-aminomethyl-4-bromo-phenyl; 2-aminosulfonyl-4-bromo-phenyl; 2-amido-4-bromo-phenyl;
 4-bromo-2-methylsulfonyl-phenyl; 2-aminomethyl-4-methyl-phenyl;
 2-aminosulfonyl-4-methyl-phenyl; 2-amido-4-methyl-phenyl; 2-methylsulfonyl-4-methyl-phenyl;
 10
 4-fluoro-pyrid-2-yl; 4-bromo-pyrid-2-yl; 4-methyl-pyrid-2-yl; 5-fluoro-thien-2-yl;
 5-bromo-thien-2-yl; 5-methyl-thien-2-yl; 2-amido-4-methoxy-phenyl;









- G_1 is absent or is selected from CH_2 , CH_2CH_2 , $\text{CH}=\text{CH}$, CH_2O , OCH_2 , NH , CH_2NH , NHCH_2 , $\text{CH}_2\text{C}(\text{O})$, $\text{C}(\text{O})\text{CH}_2$, $\text{C}(\text{O})\text{NH}$, $\text{NHC}(\text{O})$, $\text{NHC}(\text{O})\text{NH}$, $\text{C}(\text{O})\text{NHS}(\text{O})_2$, NHCOCONH , $\text{NHCOC}(\text{S})\text{NH}$, $\text{NHC}(\text{S})\text{CONH}$, $\text{CH}_2\text{S}(\text{O})_2$, $\text{S}(\text{O})_2(\text{CH}_2)$, SO_2NH , and NHSO_2 , provided that G_1 does not form a N-S, NCH_2N , NCH_2O , or NCH_2S bond with either group to which it is attached;
- 10 A is selected from cyclohexyl, indolinyl, piperidinyl, phenyl, pyridyl, and pyrimidyl, and is substituted with 0-2 R^4 ;
- 15 X is selected from CH_2 , $\text{C}(\text{O})$, $-\text{S}(\text{O})_2-$, $-\text{NHC}(\text{O})-$, $-\text{C}(\text{O})\text{NH}-$, $-\text{CH}_2\text{NH}-$, O, and $-\text{CH}_2\text{O}-$;
- 20 Y is selected from $\text{C}(\text{CH}_3)_2$, $\text{C}(\text{CH}_2\text{CH}_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentanonyl, cyclohexyl, cyclohexanonyl, pyrrolidinyl, pyrrolidinonyl, piperidinyl, piperidinonyl, tetrahydrofuranyl, and tetrahydropyranyl, and, when Y is a ring, Y is substituted with 0-1 R^4 ;
- 25 R^{1a} , at each occurrence, is selected from H, R^{1b} , $\text{CH}(\text{CH}_3)\text{R}^{1b}$, $\text{C}(\text{CH}_3)_2\text{R}^{1b}$, and CH_2R^{1b} , provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

- R^{1b} is selected from CH_3 , CH_2CH_3 , F, Cl, Br, -CN, CF_3 , OR^2 , NR^2R^{2a} , $C(O)R^{2b}$, CO_2R^{2b} , CO_2R^{2a} , $S(O)_pR^2$, $C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $NR^2SO_2R^2$, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 5-6 membered aromatic heterocycle consisting of carbon atoms and from 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-2 R^{4b} , provided that R^{1b} forms other than an O-O, N-halo, N-S, or N-CN bond;
- 10 R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, phenyl substituted with 0-1 R^{4b} , benzyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 15 R^{2a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, cyclopropyl, benzyl, phenyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 20 alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $S(O)_p$;
- 25 R^{2b} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, C_{1-5} alkyl substituted with 0-3 R^{4b} , benzyl, C_{3-6} carbocycle substituted with 0-2 R^{4b} , and 4-6 membered aromatic heterocycle substituted with

0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

5 R^{2c} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, $CH(CH_3)_2$, benzyl, phenyl substituted with 0-1 R^{4b} , and 5-6 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of carbon atoms and from 1-4
10 heteroatoms selected from the group consisting of N, O, and $S(O)_p$;

R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} carbocycle substituted with 0-2 R^{4c} , $-(CH_2)-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4
15 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, and $-(CH_2)-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p-S(O)_p$, S-O, O-N, O-S, or O-O moiety;
20

25 R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} carbocycle substituted with 0-2 R^{4c} , $-(CH_2)-C_{3-6}$ carbocycle substituted with 0-2 R^{4c} , 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4
30 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, and $-(CH_2)-5-6$ membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group

consisting of N, O, and S(O)_p, provided that R^{2e} forms other than a C(O)-halo or C(O)-S(O)_p moiety;

- R⁴, at each occurrence, is selected from OH, OR², CH₂OR²,
 5 (CH₂)₂OR², F, Br, Cl, I, CH₃, CH₂CH₃, CH₂CH₂CH₃,
 CH(CH₃)₂, CH₂CH₂CH₂CH₃, CH₂CH(CH₃)₂, CH(CH₃)CH₂CH₃,
 C(CH₃)₃, NR²R^{2a}, CH₂NR²R^{2a}, (CH₂)₂NR²R^{2a}, CF₃, and
 CF₂CF₃;
- 10 R^{4a} is selected from -(CR³R^{3g})_r-5-6 membered carbocycle
 substituted with 0-3 R^{4c}, -(CR³R^{3g})_r-5-6 membered
 heterocycle substituted with 0-3 R^{4c} and consisting of:
 carbon atoms and 1-4 heteroatoms selected from the
 group consisting of N, O, and S(O)_p, (CR³R^{3g})_rNR^{2d}R^{2d},
 15 (CR³R^{3g})_rN(→O)R^{2d}R^{2d}, (CR³R^{3g})_rOR^{2d},
 (CR³R^{3g})_r-C(O)NR^{2d}R^{2d}, (CR³R^{3g})_r-NR^{2d}C(O)R^{2e},
 (CR³R^{3g})_r-C(O)R^{2e}, (CR³R^{3g})_r-NR^{2d}C(O)NR^{2d}R^{2d},
 (CR³R^{3g})_r-NR^{2d}C(O)OR^{2d}, (CR³R^{3g})_r-NR^{2d}SO₂R^{2d}, and
 (CR³R^{3g})_r-S(O)_pR^{2d}, provided that S(O)_pR^{2d} forms other
 20 than S(O)₂H or S(O)H;

- R^{4b}, at each occurrence, is selected from H, =O, OR³,
 CH₂OR³, F, Cl, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, -CN,
 NO₂, NR³R^{3a}, CH₂NR³R^{3a}, C(O)R³, C(O)OR^{3c}, NR³C(O)R^{3a},
 25 C(O)NR³R^{3a}, SO₂NR³R^{3a}, NR³SO₂-C₁₋₄ alkyl, NR³SO₂-phenyl,
 S(O)_p-C₁₋₄ alkyl, S(O)_p-phenyl, and CF₃;

- R^{4c}, at each occurrence, is selected from =O, OR², CH₂OR²,
 F, Br, Cl, CF₃, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, C₂₋₃
 30 alkenyl, C₂₋₃ alkynyl, -CN, NO₂, NR²R^{2a}, CH₂NR²R^{2a},
 N(→O)R²R^{2a}, CH₂N(→O)R²R^{2a}, C(O)R^{2c}, CH₂C(O)R^{2c},
 NR²C(O)R^{2b}, CH₂NR²C(O)R^{2b}, C(O)NR²R^{2a}, CH₂C(O)NR²R^{2a},

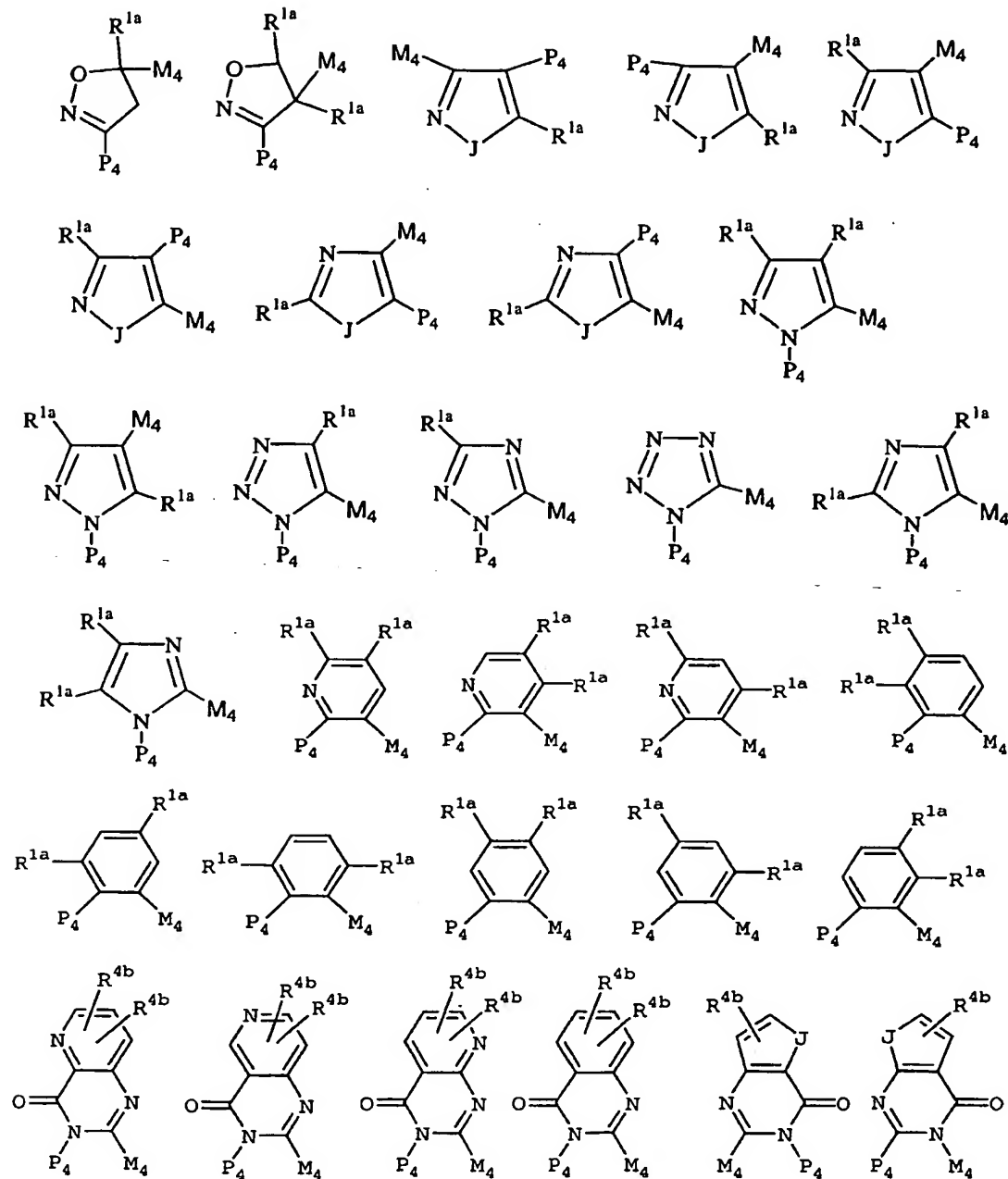
$\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{CH}_2\text{SO}_2\text{NR}^2\text{R}^{2a}$, $\text{NR}^2\text{SO}_2\text{R}^{5a}$, $\text{CH}_2\text{NR}^2\text{SO}_2\text{R}^{5a}$,
 $\text{S(O)}_p\text{R}^{5a}$, $\text{CH}_2\text{S(O)}_p\text{R}^{5a}$, CF_3 , CF_2CF_3 , C_{3-6} carbocycle
substituted with 0-2 R^{4b} , $(\text{CH}_2)_{\text{C}_{3-6}}$ carbocycle
substituted with 0-2 R^{4b} , 5-6 membered heterocycle
5 substituted with 0-2 R^{4b} and consisting of carbon atoms
and from 1-4 heteroatoms selected from the group
consisting of N, O, and S(O)_p , and $(\text{CH}_2)_{5-6}$ membered
heterocycle substituted with 0-2 R^{4b} and consisting of
carbon atoms and from 1-4 heteroatoms selected from
10 the group consisting of N, O, and S(O)_p ;

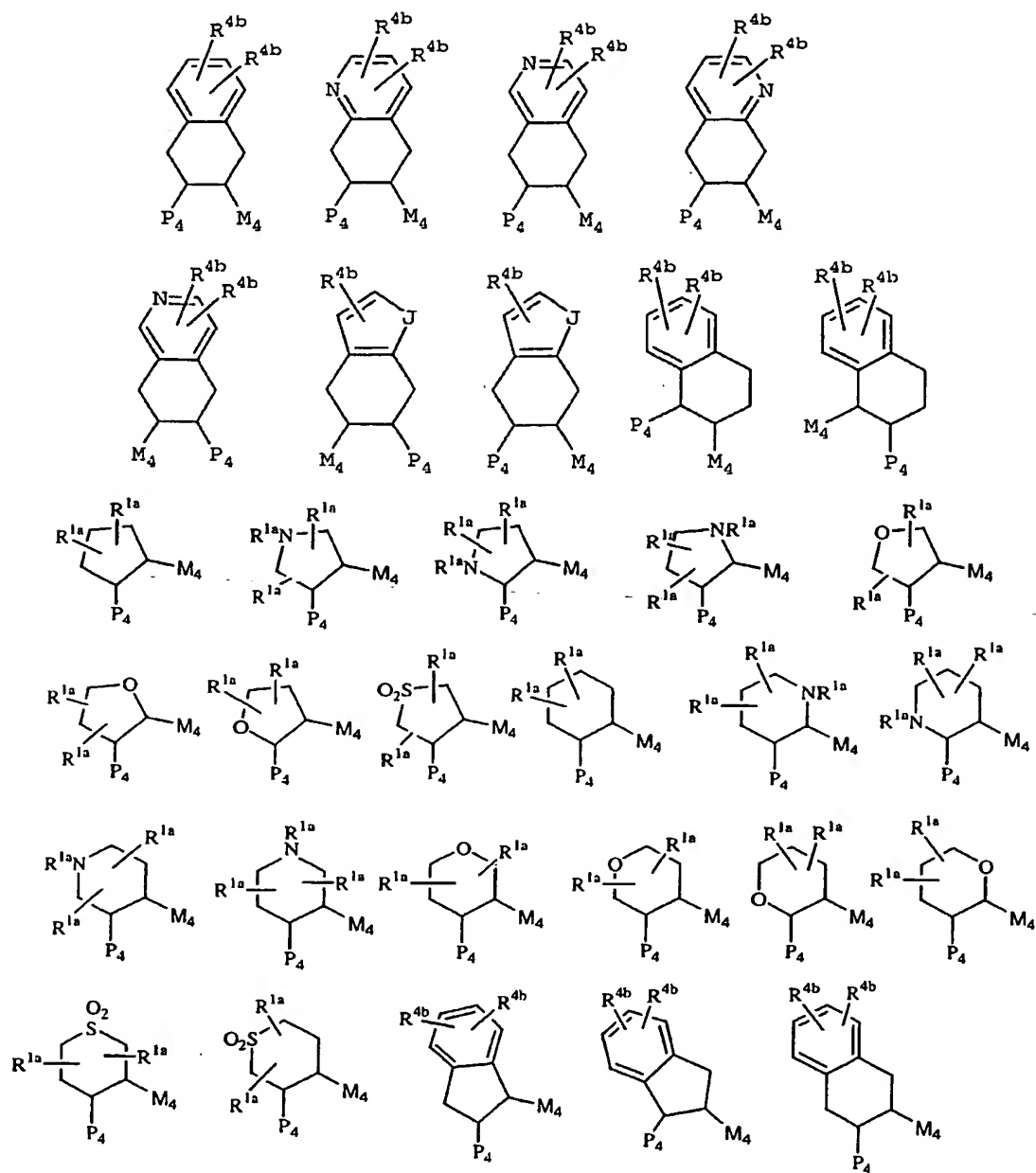
R^5 , at each occurrence, is selected from H, =O, CH_3 , CH_2CH_3 ,
 $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, OR^3 , CH_2OR^3 , F, Cl, -CN, NO_2 ,
 NR^3R^{3a} , $\text{CH}_2\text{NR}^3\text{R}^{3a}$, $\text{C(O)}\text{R}^3$, $\text{C(O)}\text{OR}^{3c}$, $\text{NR}^3\text{C(O)}\text{R}^{3a}$,
15 $\text{C(O)}\text{NR}^3\text{R}^{3a}$, $\text{SO}_2\text{NR}^3\text{R}^{3a}$, $\text{NR}^3\text{SO}_2\text{-C}_{1-4}$ alkyl, $\text{NR}^3\text{SO}_2\text{-phenyl}$,
 $\text{S(O)}_p\text{-C}_{1-4}$ alkyl, $\text{S(O)}_p\text{-phenyl}$, CF_3 , phenyl substituted
with 0-2 R^6 , naphthyl substituted with 0-2 R^6 , and
benzyl substituted with 0-2 R^6 ; and,

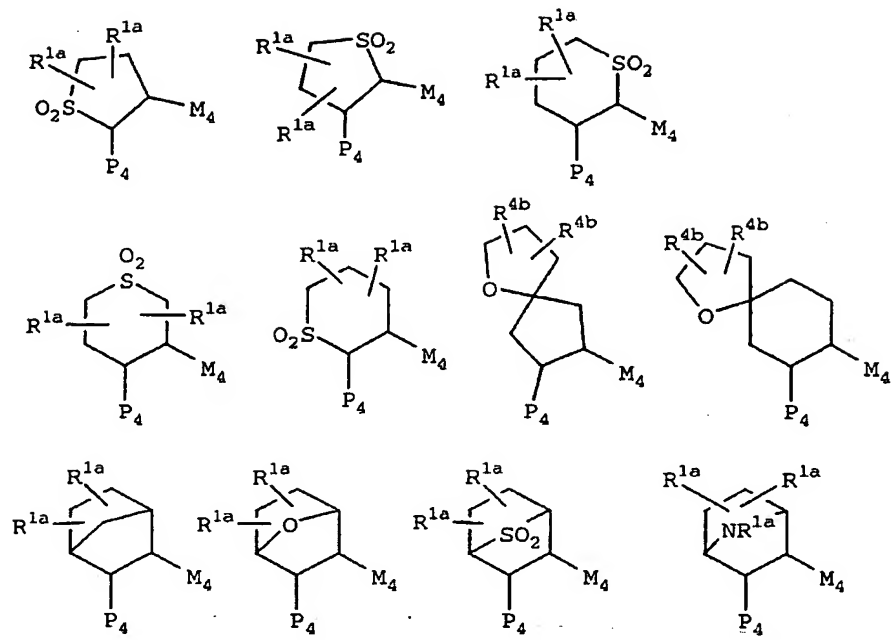
20 R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl,
 CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, -CN, NO_2 , NR^2R^{2a} ,
 $\text{CH}_2\text{NR}^2\text{R}^{2a}$, $\text{C(O)}\text{R}^{2b}$, $\text{CH}_2\text{C(O)}\text{R}^{2b}$, $\text{NR}^2\text{C(O)}\text{R}^{2b}$, and
 $\text{SO}_2\text{NR}^2\text{R}^{2a}$.

25

12. A compound according to Claim 11, wherein the
compound is selected from:







J is selected from O, S, NH, and NR^{1a} ;

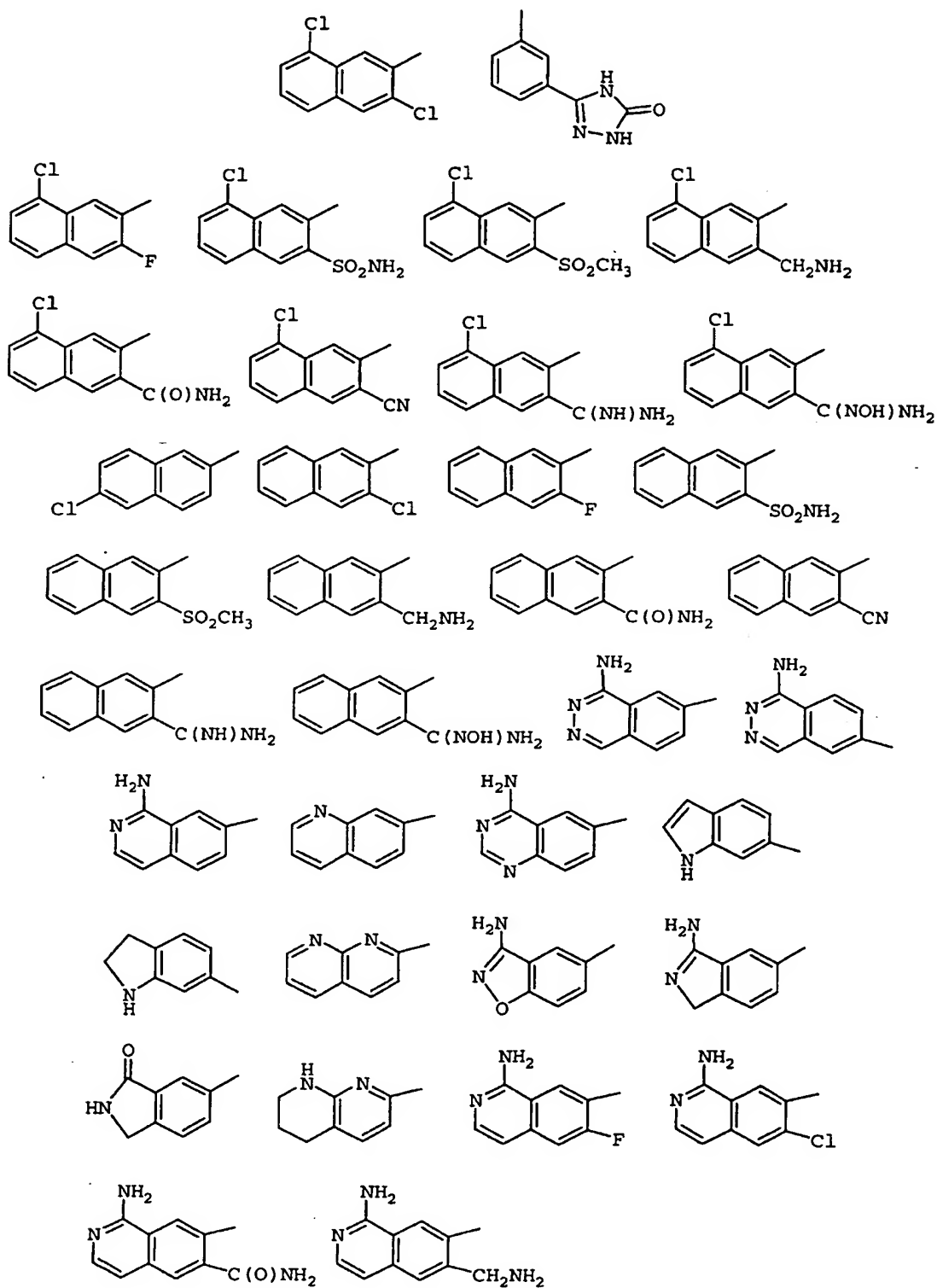
5 P_4 is $-G_1-G$;

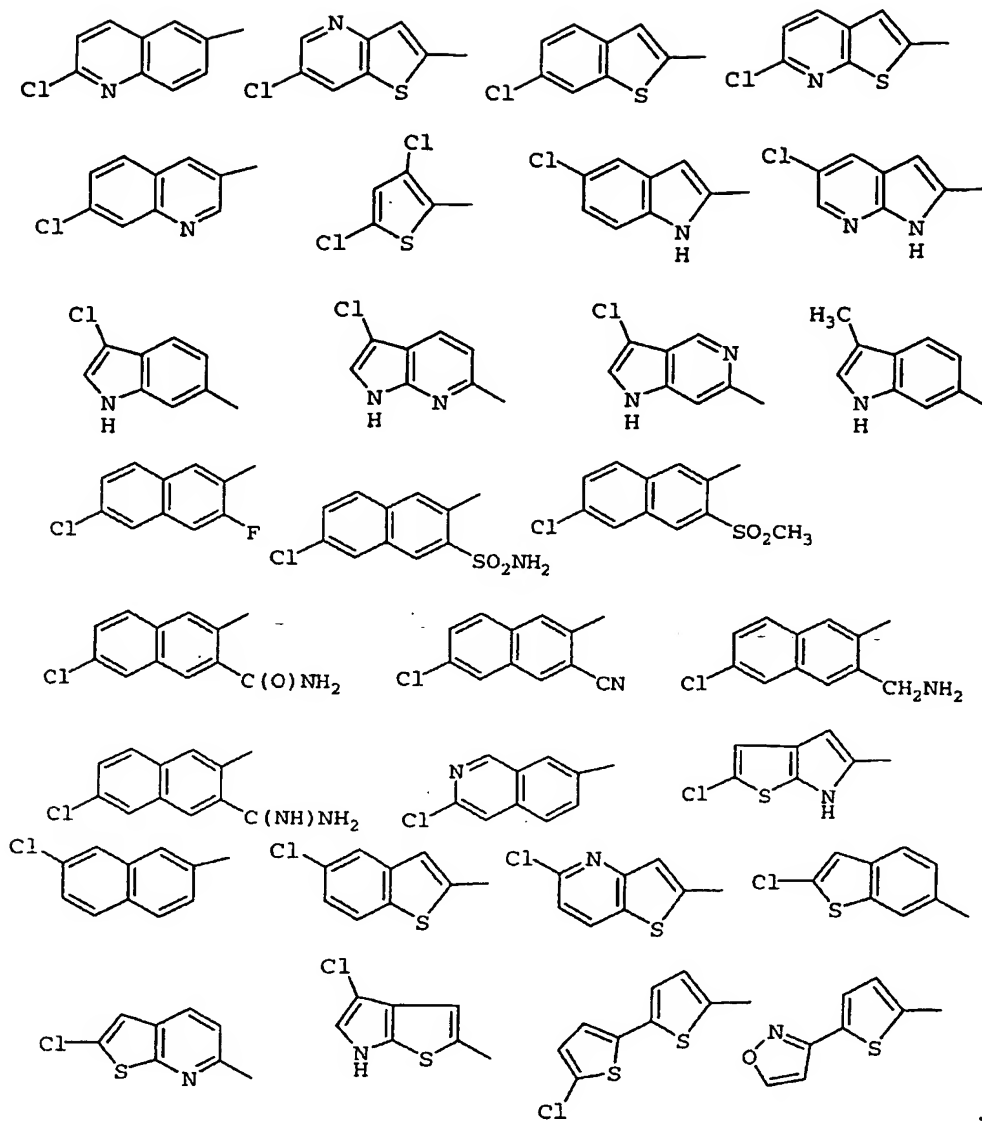
M_4 is $-Z-A-B$;

G is selected from:

- 10 2-amido-4-methoxy-phenyl; 2-amido-phenyl;
2-aminomethyl-3-fluoro-phenyl;
2-aminomethyl-4-fluoro-phenyl;
2-aminomethyl-5-fluoro-phenyl;
2-aminomethyl-6-fluoro-phenyl; 2-aminomethyl-phenyl;
- 15 2-amino-pyrid-4-yl; 2-aminosulfonyl-4-methoxy-phenyl;
2-aminosulfonyl-phenyl; 3-amido-phenyl;
3-amino-4-chloro-phenyl; 3-aminomethyl-phenyl;
3-chloro-phenyl; 4-chloro-phenyl; 4-methoxy-phenyl;
5-chloro-pyrid-2-yl; 5-chloro-thien-2-yl;
- 20 6-amino-5-chloro-pyrid-2-yl; 6-amino-pyrid-2-yl; 5-chloro-pyrimidin-3-yl; 6-chloro-pyridazin-3-yl;
2-aminomethyl-4-chloro-phenyl;

2-aminosulfonyl-4-chloro-phenyl; 2-amido-4-chloro-phenyl;
4-chloro-2-methylsulfonyl-phenyl;





- 5 G_1 is absent or is selected from CH=CH, CH₂NH, NHCH₂, CH₂C(O), C(O)CH₂, C(O)NH, NHC(O), NHC(O)NH, CH₂S(O)₂, S(O)₂(CH₂), SO₂NH, and NHSO₂, provided that G_1 does not form a N-S, NCH₂N, NCH₂O, or NCH₂S bond with either group to which it is attached;

10

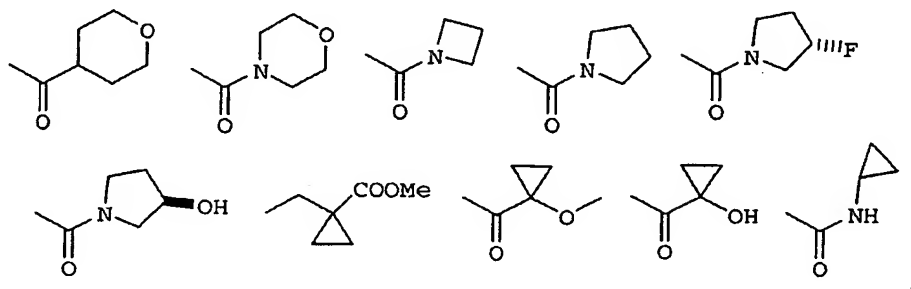
A is selected from the group: cyclohexyl, indolinyl, piperidinyl, phenyl, 2-pyridyl, 3-pyridyl, 2-pyrimidyl, 2-Cl-phenyl, 3-Cl-phenyl, 2-F-phenyl, 3-F-

phenyl, 2-methylphenyl, 2-aminophenyl, and 2-methoxyphenyl;

- Y is selected from $C(CH_3)_2$, $C(CH_2CH_3)_2$, cyclopropyl, cyclobutyl, cyclopentyl, 2-cyclopentanonyl, cyclohexyl, 2-cyclohexanonyl, pyrrolidiny (attached to A and R^{4a} at the 2-position), pyrrolidiny (attached to A and R^{4a} at the 3-position), 2-pyrrolidinonyl (attached to A and R^{4a} at the 3-position), piperidiny (attached to A and R^{4a} at the 4-position), 4-piperidinonyl (attached to A and R^{4a} at the 3-position), tetrahydrofurany, and tetrahydropyrany (attached to A and R^{4a} at the 4-position);
- R^{1a} , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$, CH_2F , CH_2Cl , Br, CH_2Br , -CN, CH_2CN , CF_3 , CH_2CF_3 , OCH_3 , CH_2OH , $C(CH_3)_2OH$, CH_2OCH_3 , $CH_2CH_2OCH_3$, NH_2 , CH_2NH_2 , $NHCH_3$, CH_2NHCH_3 , $N(CH_3)_2$, $CH_2N(CH_3)_2$, CO_2H , CH_2CO_2H , $CH_2CH_2CO_2H$, $COCH_3$, CO_2CH_3 , $CH_2CO_2CH_3$, SCH_3 , CH_2SCH_3 , $S(O)CH_3$, $CH_2S(O)CH_3$, $S(O)_2CH_3$, $CH_2S(O)_2CH_3$, $C(O)NH_2$, $CH_2C(O)NH_2$, SO_2NH_2 , $CH_2SO_2NH_2$, $NHSO_2CH_3$, $CH_2NHSO_2CH_3$, $COCH_2C(CH_3)_3$, $COCH_2OH$, $COCH_2OCH_3$, $COC(CH_3)_2OH$, $COC(CH_3)_2CH_2OH$, $COC(CH_3)_2CH_2OCH_3$, $C(O)OCH_2CH_2OCH_3$, $COCF_3$, $CO_2CH_2CH_3$, $CO_2CH(CH_3)_2$, $CO_2C(CH_3)_3$, $CH_2CH_2CO_2CH_2CH_3$, $CONH(CH_3)$, $CONH(CH_2CH_3)$, $CONHC(CH_3)_3$, $CON(CH_3)_2$, $CON(CH_3)(CH_2CH_3)$, $CON(CH_3)CH(CH_3)_2$, $CH_2CON(CH_3)_2$, C(O)-phenyl, C(O)-cyclopropyl, C(O)-cyclobutyl, C(O)-cyclopentyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyridin-2-yl-N-oxide, pyridin-3-yl-N-oxide, pyridin-4-yl-N-oxide, imidazol-1-yl, CH_2 -imidazol-1-yl, 4-methyloxazol-2-yl, 4-N,N-dimethylaminomethyl-oxazol-2-yl, 1,2,3,4-tetrazol-1-yl, 1,2,3,4-tetrazol-5-yl, CH_2 -

1,2,3,4-tetrazol-1-yl, and CH_2 -1,2,3,4-tetrazol-5-yl, provided that R^{1a} forms other than an N-halo, N-S, or N-CN bond;

5 alternatively, R^{1a} is selected from:



- R^2 , at each occurrence, is selected from H, CH_3 , CH_2CH_3 , $\text{CH}_2\text{CH}_2\text{CH}_3$, $\text{CH}(\text{CH}_3)_2$, phenyl substituted with 0-1 R^{4b} , benzyl substituted with 0-1 R^{4b} , and 5 membered aromatic heterocycle substituted with 0-1 R^{4b} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $\text{S}(\text{O})_p$;
- 10
- 15 R^{2a} , at each occurrence, is selected from H, CH_3 , and CH_2CH_3 ;

alternatively, R^2 and R^{2a} , together with the nitrogen atom to which they are attached, combine to form a 3-6 membered saturated, partially saturated or unsaturated ring substituted with 0-1 R^{4b} and consisting of: 0-1 additional heteroatoms selected from the group consisting of N, O, and $\text{S}(\text{O})_p$;

20

25 R^{2b} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , CH_3 , and CH_2CH_3 ;

R^{2c} , at each occurrence, is selected from OH, OCH_3 , OCH_2CH_3 , CH_3 , and CH_2CH_3 ;

R^{2d} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c} consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p$ - $S(O)_p$, S-O, O-N, O-S, or O-O moiety;

R^{2e} , at each occurrence, is selected from H, R^{4c} , C_{1-4} alkyl substituted with 0-2 R^{4c} , C_{3-6} cycloalkyl substituted with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a C(O)-halo or C(O)- $S(O)_p$ moiety;

R^{4a} is selected from $-(CH_2)_r$ -5-6 membered carbocycle substituted with 0-3 R^{4c} , $-(CH_2)_r$ -5-6 membered heterocycle substituted with 0-3 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, $(CH_2)_rNR^{2d}R^{2d}$, $(CH_2)_rN(\rightarrow O)R^{2d}R^{2d}$, $(CH_2)_rOR^{2d}$, $(CH_2)_r-C(O)NR^{2d}R^{2d}$, $(CH_2)_r-NR^{2d}C(O)R^{2e}$, $(CH_2)_r-C(O)R^{2e}$, $(CH_2)_r-NR^{2d}C(O)NR^{2d}R^{2d}$, $(CH_2)_r-NR^{2d}C(O)OR^{2d}$, $(CH_2)_r-NR^{2d}SO_2R^{2d}$, and $(CH_2)_r-S(O)_pR^{2d}$, provided that $S(O)_pR^{2d}$ forms other than $S(O)_2H$ or $S(O)H$;

R^{4b} , at each occurrence, is selected from H, $=O$, OR^3 , CH_2OR^3 , F, Cl, CH_3 , CH_2CH_3 , NR^3R^{3a} , $CH_2NR^3R^{3a}$, $C(O)R^3$,

$C(O)OR^3c$, $NR^3C(O)R^3a$, $C(O)NR^3R^3a$, $SO_2NR^3R^3a$,
 NR^3SO_2 -phenyl, $S(O)_2CH_3$, $S(O)_2$ -phenyl, and CF_3 ;

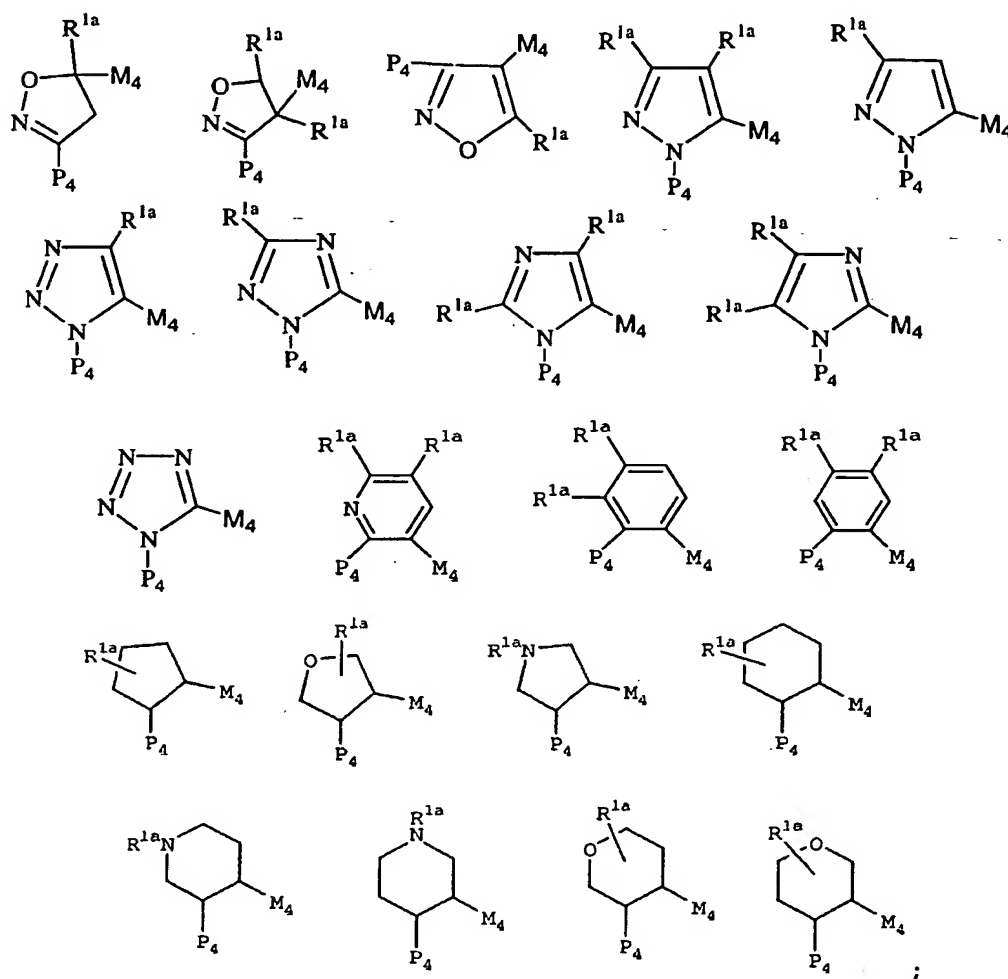
R^{4c} , at each occurrence, is selected from $=O$, OH , OCH_3 ,
 5 OCH_2CH_3 , $OCH_2CH_2CH_3$, $OCH(CH_3)_2$, CH_3 , CH_2CH_3 , $CH_2CH_2CH_3$,
 $CH(CH_3)_2$, C_{2-3} alkenyl, C_{2-3} alkynyl, CH_2OH , CH_2OCH_3 ,
 $CH_2OCH_2CH_3$, $CH_2OCH_2CH_2CH_3$, $CH_2OCH(CH_3)_2$, F , Br , Cl , CF_3 ,
 NR^2R^{2a} , $CH_2NR^2R^{2a}$, $N(\rightarrow O)R^2R^{2a}$, $CH_2N(\rightarrow O)R^2R^{2a}$, $C(O)R^{2c}$,
 $CH_2C(O)R^{2c}$, $NR^2C(O)R^{2b}$, $CH_2NR^2C(O)R^{2b}$, $C(O)NR^2R^{2a}$,
 10 $CH_2C(O)NR^2R^{2a}$, $SO_2NR^2R^{2a}$, $CH_2SO_2NR^2R^{2a}$, $NR^2SO_2R^{5a}$,
 $CH_2NR^2SO_2R^{5a}$, $S(O)_pR^{5a}$, $CH_2S(O)_pR^{5a}$, CF_3 , cyclopropyl
 substituted with 0-1 R^{4b} , cyclobutyl substituted with
 0-1 R^{4b} , cyclopentyl substituted with 0-1 R^{4b} , phenyl
 substituted with 0-1 R^{4b} , $-CH_2$ -cyclopropyl substituted
 15 with 0-1 R^{4b} , $-CH_2$ -cyclobutyl substituted with 0-1 R^{4b} ,
 $-CH_2$ -cyclopentyl substituted with 0-1 R^{4b} , benzyl
 substituted with 0-2 R^{4b} , 5-6 membered aromatic
 heterocycle substituted with 0-2 R^{4b} and consisting of
 carbon atoms and from 1-4 heteroatoms selected from
 20 the group consisting of N , O , and $S(O)_p$, and $(CH_2)_5$ -6
 membered aromatic heterocycle substituted with 0-2 R^{4b}
 and consisting of carbon atoms and from 1-4
 heteroatoms selected from the group consisting of N ,
 O , and $S(O)_p$;

25 R^5 , at each occurrence, is selected from H , $=O$, CH_3 , CH_2CH_3 ,
 OR^3 , CH_2OR^3 , F , Cl , NR^3R^3a , $CH_2NR^3R^3a$, $C(O)R^3$, $C(O)OR^3c$,
 $NR^3C(O)R^3a$, $C(O)NR^3R^3a$, $SO_2NR^3R^3a$, NR^3SO_2 - C_{1-4} alkyl,
 NR^3SO_2 -phenyl, $S(O)_2$ - CH_3 , $S(O)_2$ -phenyl, CF_3 , phenyl
 30 substituted with 0-2 R^6 , naphthyl substituted with 0-2
 R^6 , and benzyl substituted with 0-2 R^6 ; and,

R^6 , at each occurrence, is selected from H, OH, OR^2 , F, Cl, CH_3 , CH_2CH_3 , NR^2R^{2a} , $CH_2NR^2R^{2a}$, $C(O)R^{2b}$, $CH_2C(O)R^{2b}$, $NR^2C(O)R^{2b}$, and $SO_2NR^2R^{2a}$.

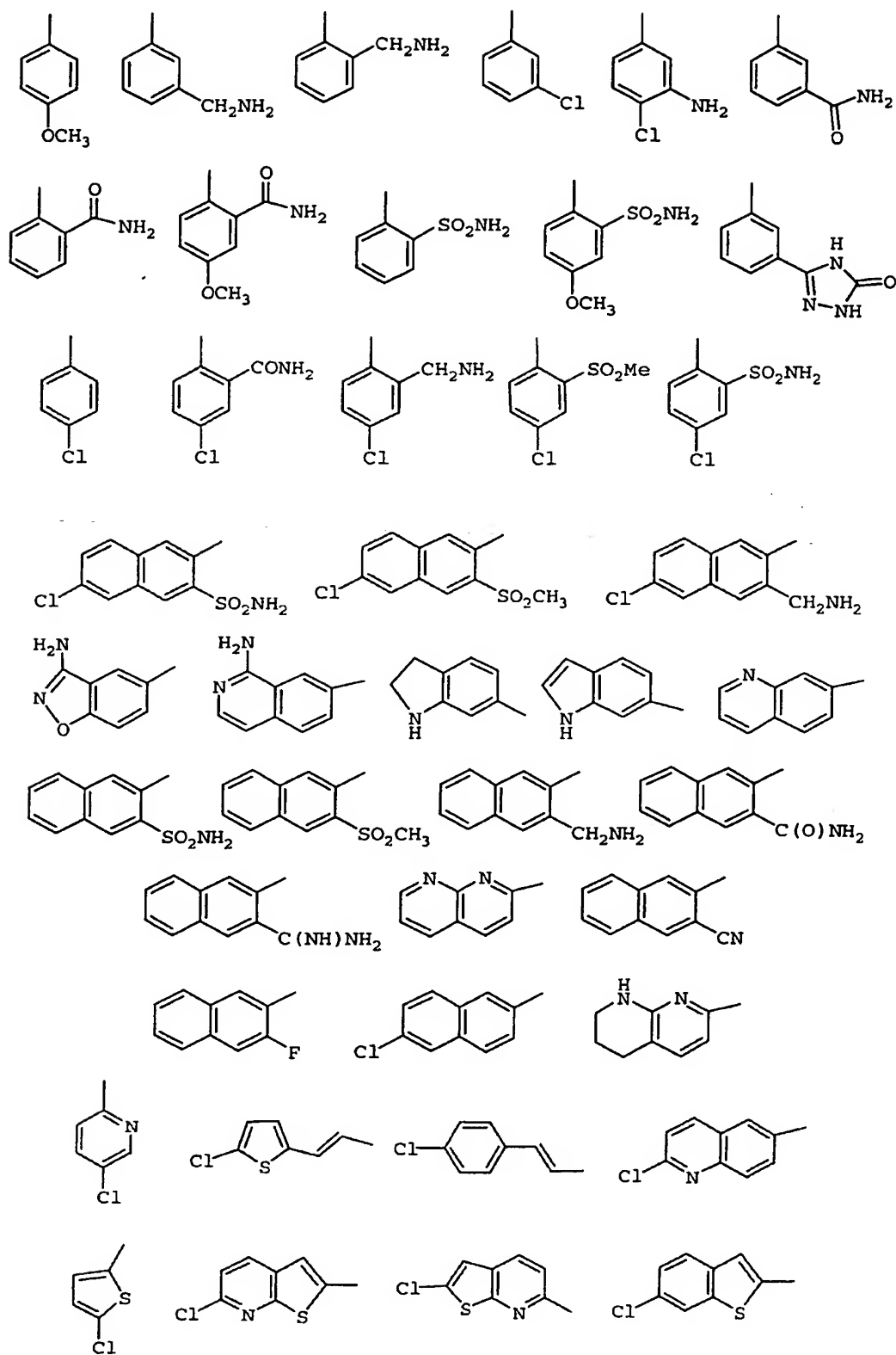
5

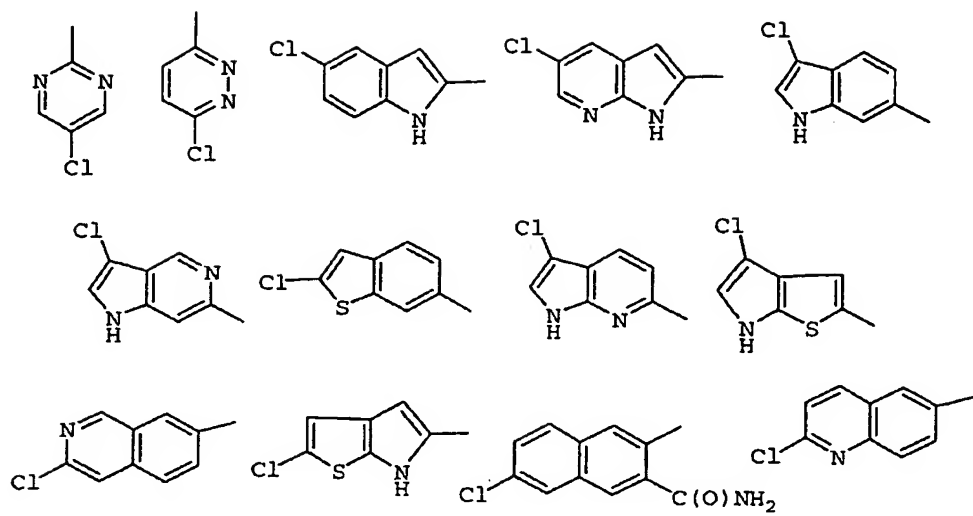
13. A compound according to Claim 12, wherein the compound is selected from:



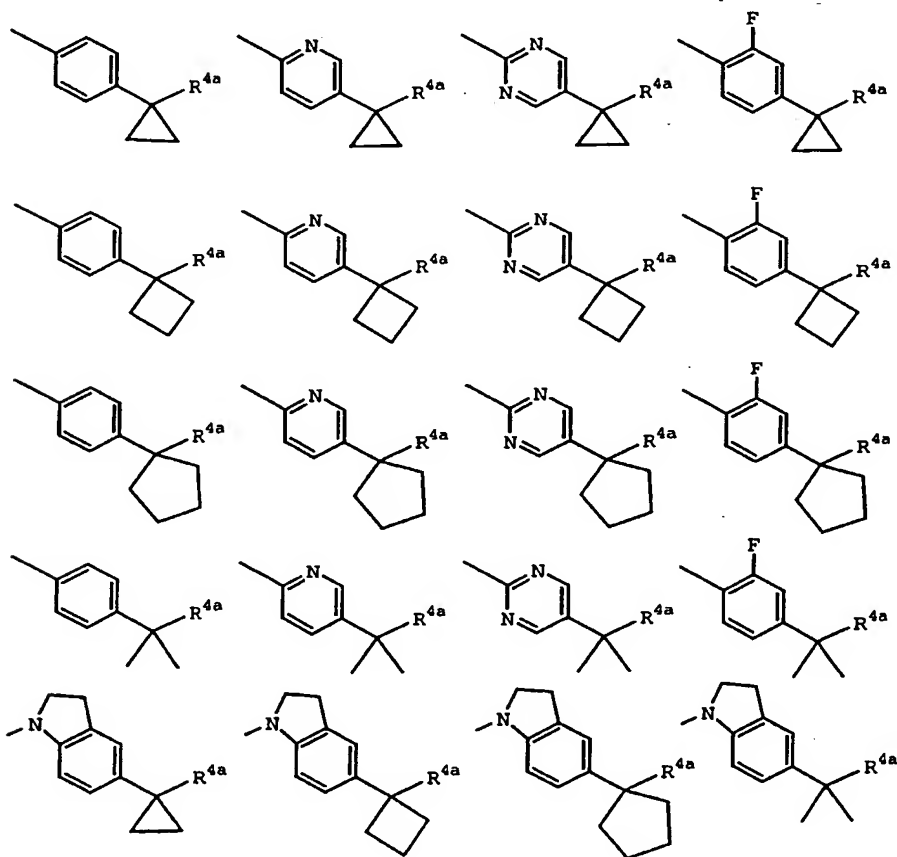
10

$-G_1-G$ is selected from:





A-B is selected from:



5

R^{2d}, at each occurrence, is selected from H, C₁₋₄ alkyl substituted with 0-1 R^{4c}, C₃₋₆ cycloalkyl substituted

with 0-2 R^{4c} , phenyl substituted with 0-2 R^{4c} , and a 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2d} forms other than a N-halo, N-C-halo, $S(O)_p$ -halo, O-halo, N-S, S-N, $S(O)_p$ - $S(O)_p$, S-O, O-N, O-S, or O-O moiety;

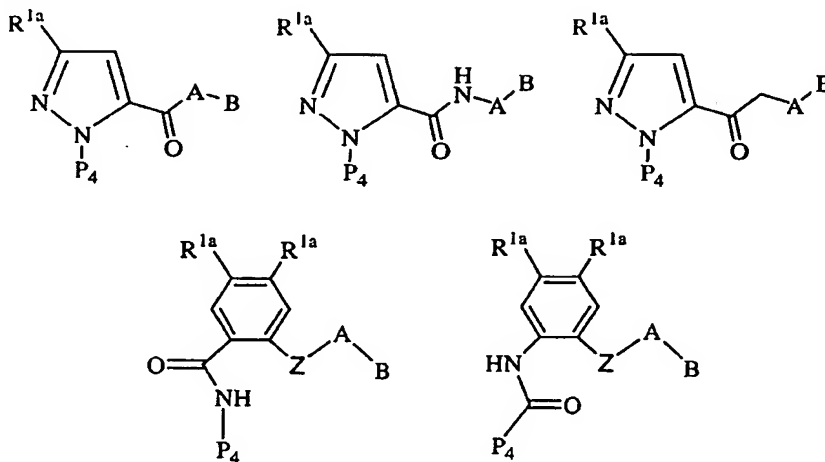
R^{2e} , at each occurrence, is selected from H, C_{1-4} alkyl substituted with 0-1 R^{4c} , C_{3-6} cycloalkyl substituted with 0-2 R^{4c} , phenyl, substituted with 0-2 R^{4c} , and 5-6 membered aromatic heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, provided that R^{2e} forms other than a $C(O)$ -halo or $C(O)$ - $S(O)_p$ moiety;

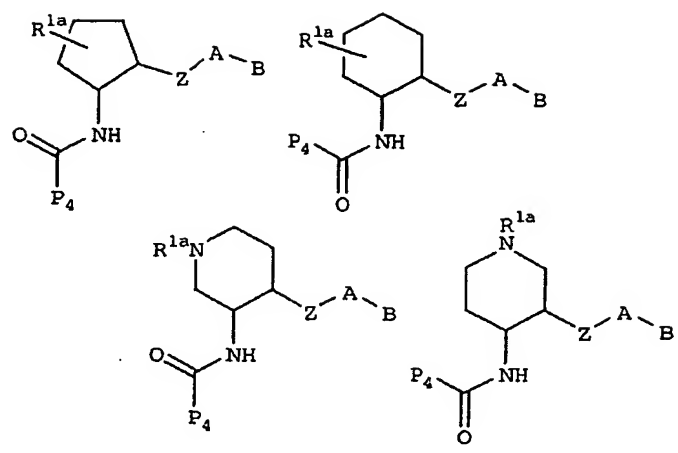
R^{4a} is selected from $NR^{2d}R^{2d}$, $CH_2NR^{2d}R^{2d}$, $CH_2CH_2NR^{2d}R^{2d}$, $N(\rightarrow O)R^{2d}R^{2d}$, $CH_2N(\rightarrow O)R^{2d}R^{2d}$, CH_2OR^{2d} , $C(O)R^{2e}$, $C(O)NR^{2d}R^{2d}$, $CH_2C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)R^{2e}$, $CH_2NR^{2d}C(O)R^{2e}$, $NR^{2d}C(O)NR^{2d}R^{2d}$, $CH_2NR^{2d}C(O)NR^{2d}R^{2d}$, $NR^{2d}C(O)OR^{2d}$, $CH_2NR^{2d}C(O)OR^{2d}$, $NR^{2d}SO_2R^{2d}$, $CH_2NR^{2d}SO_2R^{2d}$, $S(O)_pR^{2d}$, $CH_2S(O)_pR^{2d}$, 5-6 membered carbocycle substituted with 0-2 R^{4c} , $-(CH_2)$ -5-6 membered carbocycle substituted with 0-2 R^{4c} , $-(CH_2)_2$ -5-6 membered carbocycle substituted with 0-2 R^{4c} , 5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, $-(CH_2)$ -5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$, and $-(CH_2)_2$ -5-6 membered heterocycle substituted with 0-2 R^{4c} and consisting of: carbon atoms and 1-4 heteroatoms

selected from the group consisting of N, O, and S(O)_p provided that S(O)_pR^{2d} forms other than S(O)₂H or S(O)H; and,

- 5 R^{4c} is selected from =O, OH, OCH₃, OCH₂CH₃, OCH₂CH₂CH₃, OCH(CH₃)₂, CH₃, CH₂CH₃, CH₂CH₂CH₃, CH(CH₃)₂, CH=CH₂, CH≡CH, CH₂OH, CH₂OCH₃, CH₂OCH₂CH₃, CH₂OCH₂CH₂CH₃, CH₂OCH(CH₃)₂, F, Br, Cl, CF₃, NR²R^{2a}, CH₂NR²R^{2a}, C(O)R^{2c}, CH₂C(O)R^{2c}, NR²C(O)R^{2b}, CH₂NR²C(O)R^{2b},
 10 C(O)NR²R^{2a}, CH₂C(O)NR²R^{2a}, SO₂NR²R^{2a}, CH₂SO₂NR²R^{2a}, NR²SO₂R^{5a}, CH₂NR²SO₂R^{5a}, S(O)_pR^{5a}, and CH₂S(O)_pR^{5a}.

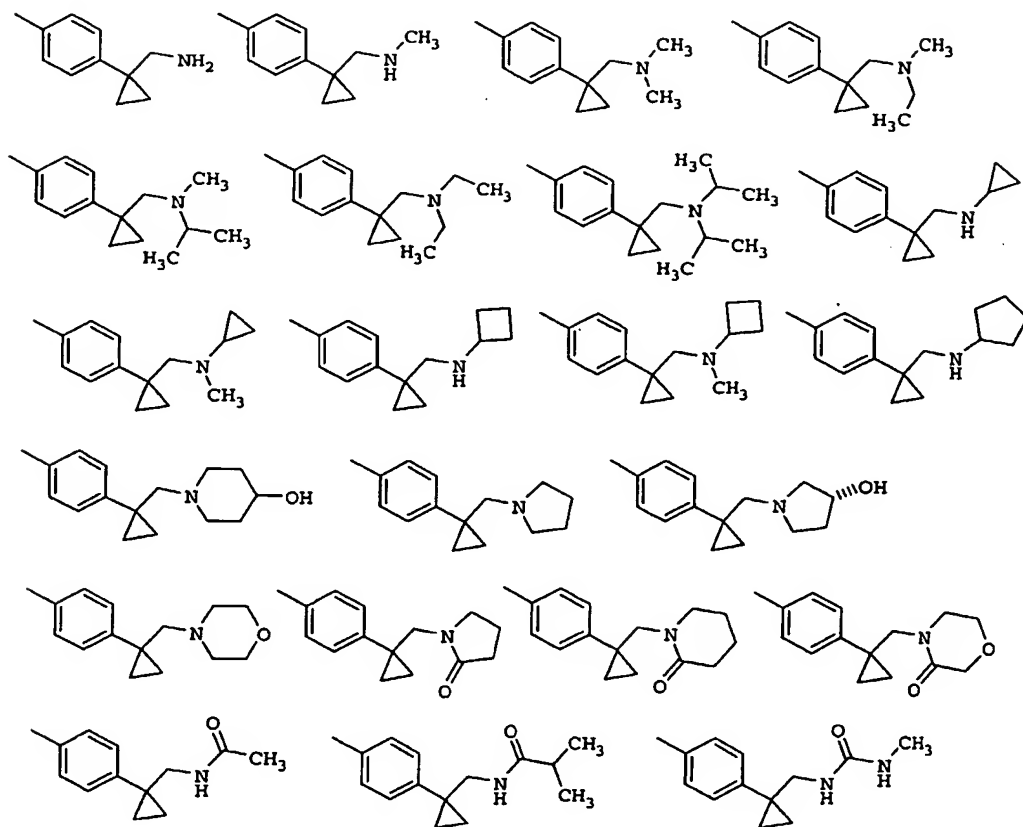
14. A compound according to Claim 13, wherein the compound
 15 is selected from:

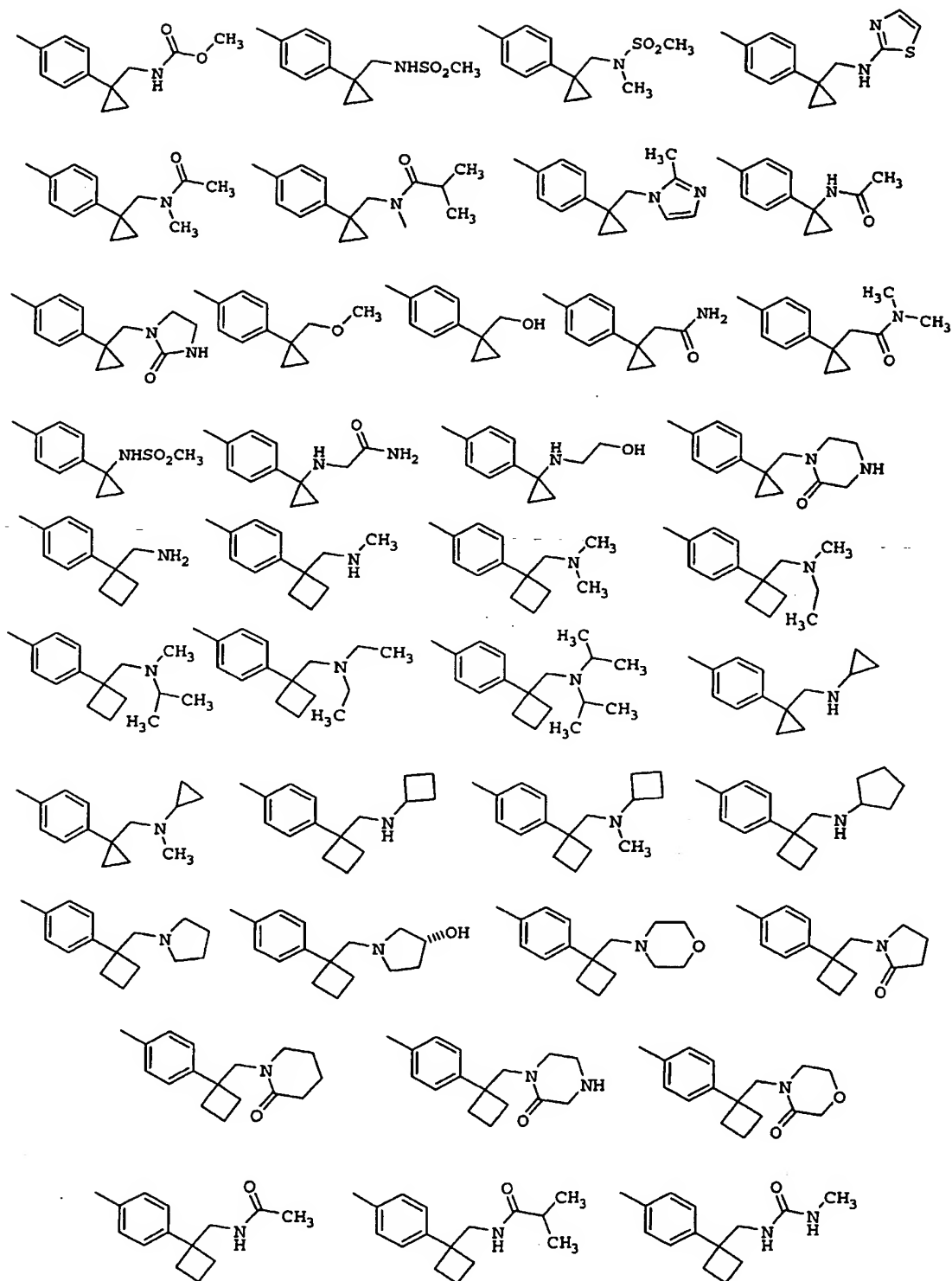


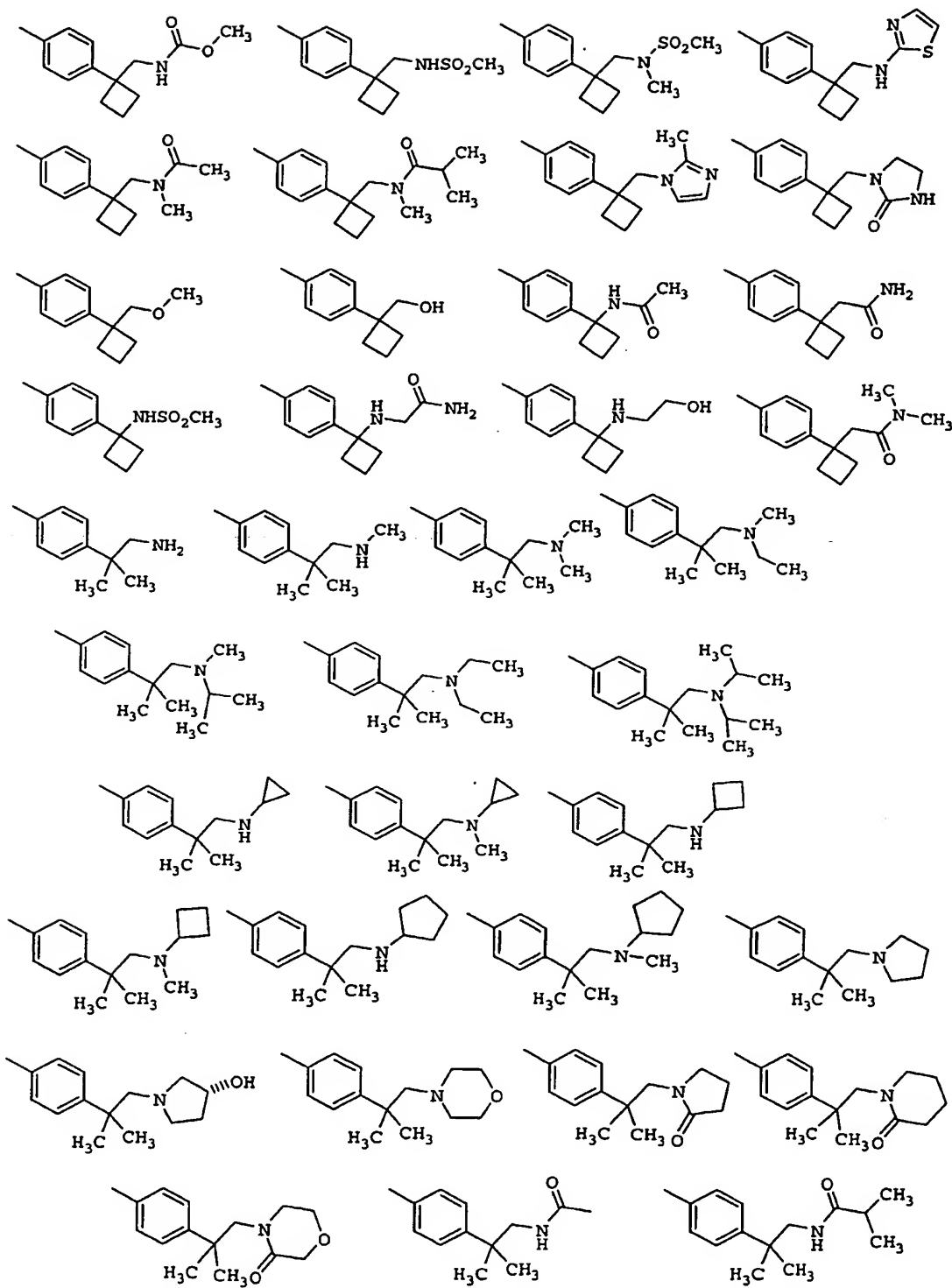


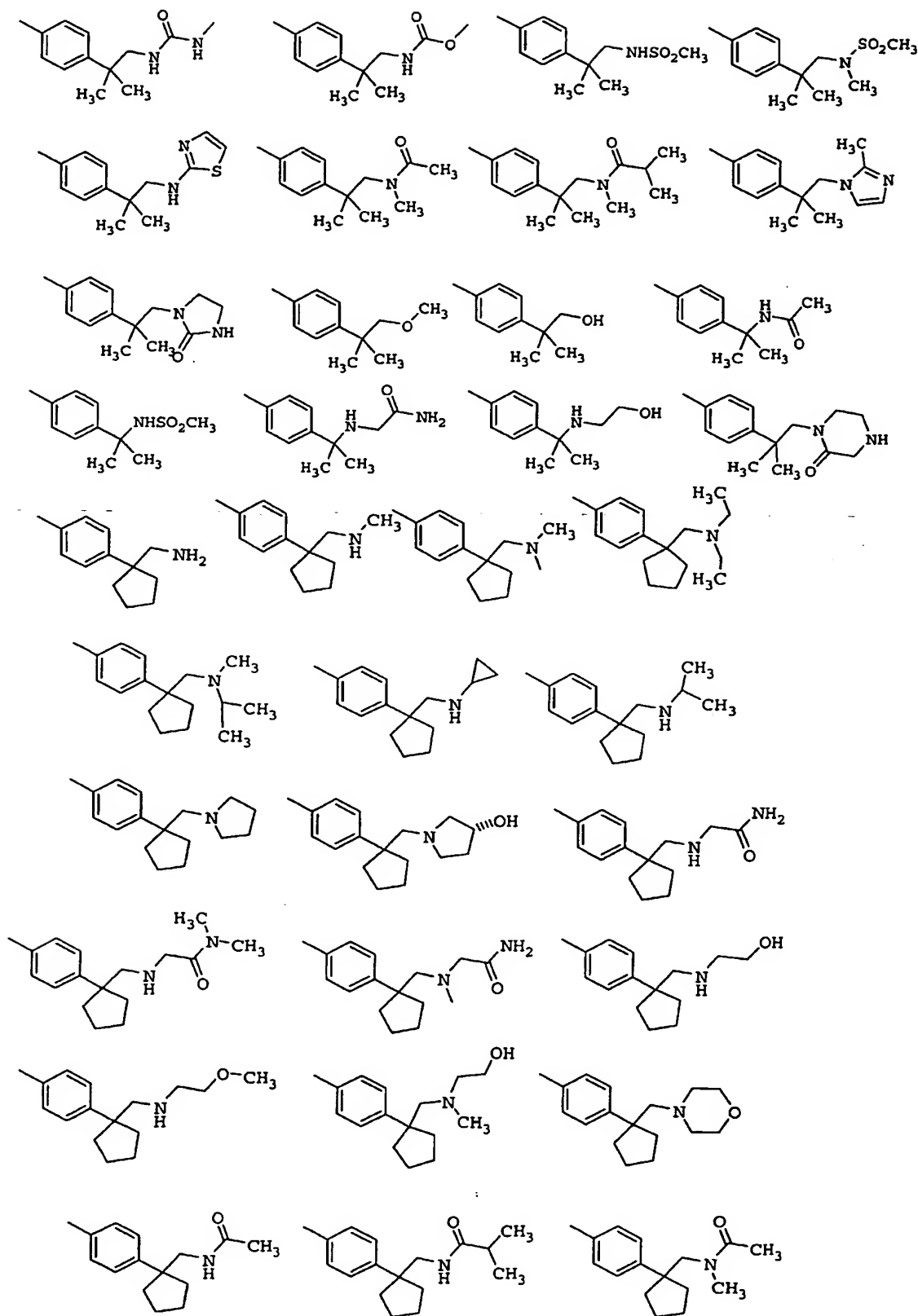
Z is selected from a $NHCH_2$, $C(O)NH$, $NHC(O)$, and $NHSO_2$; and,

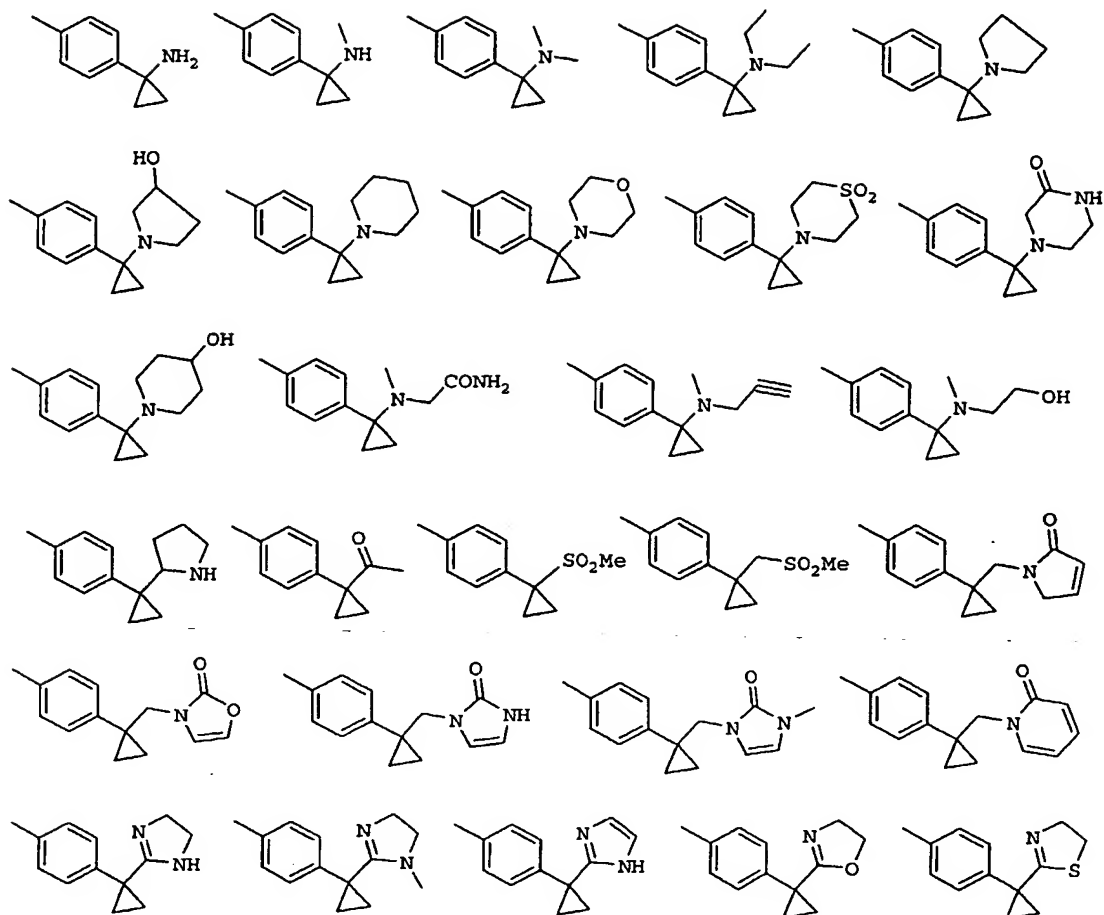
5 $A-B$ is selected from:

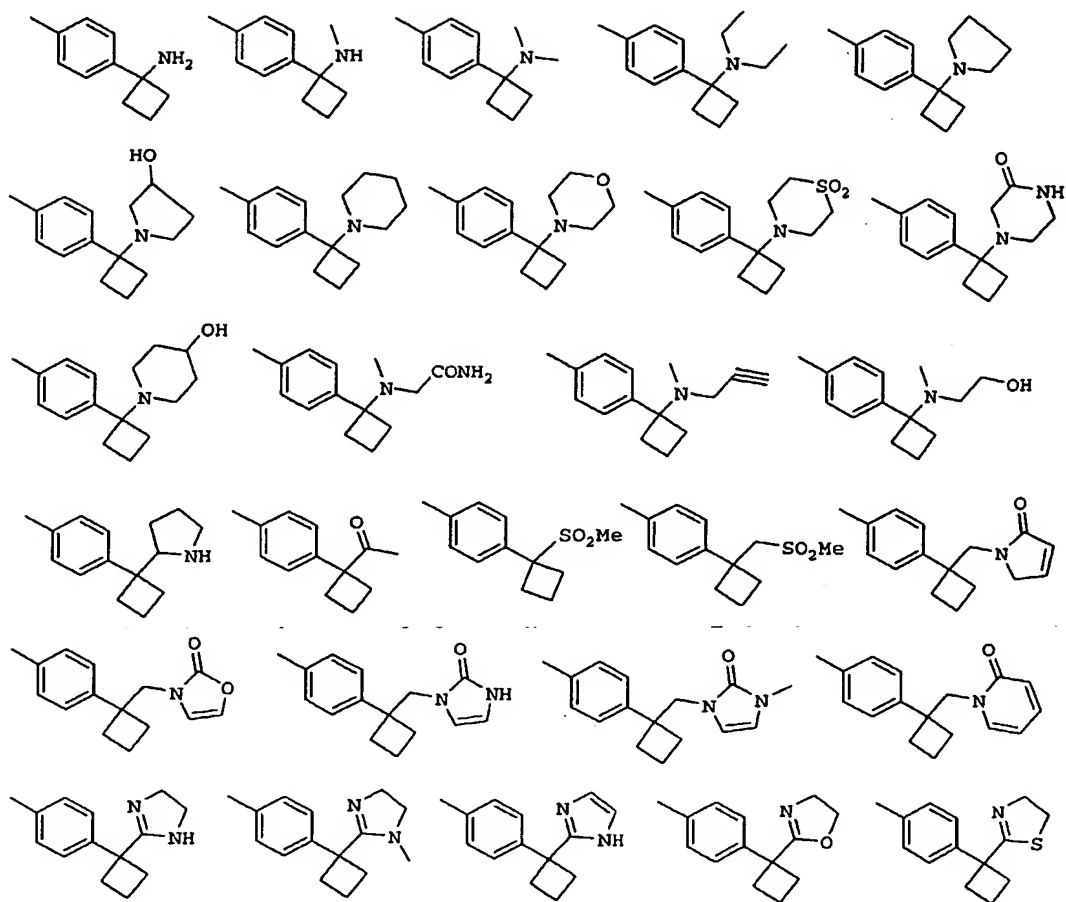


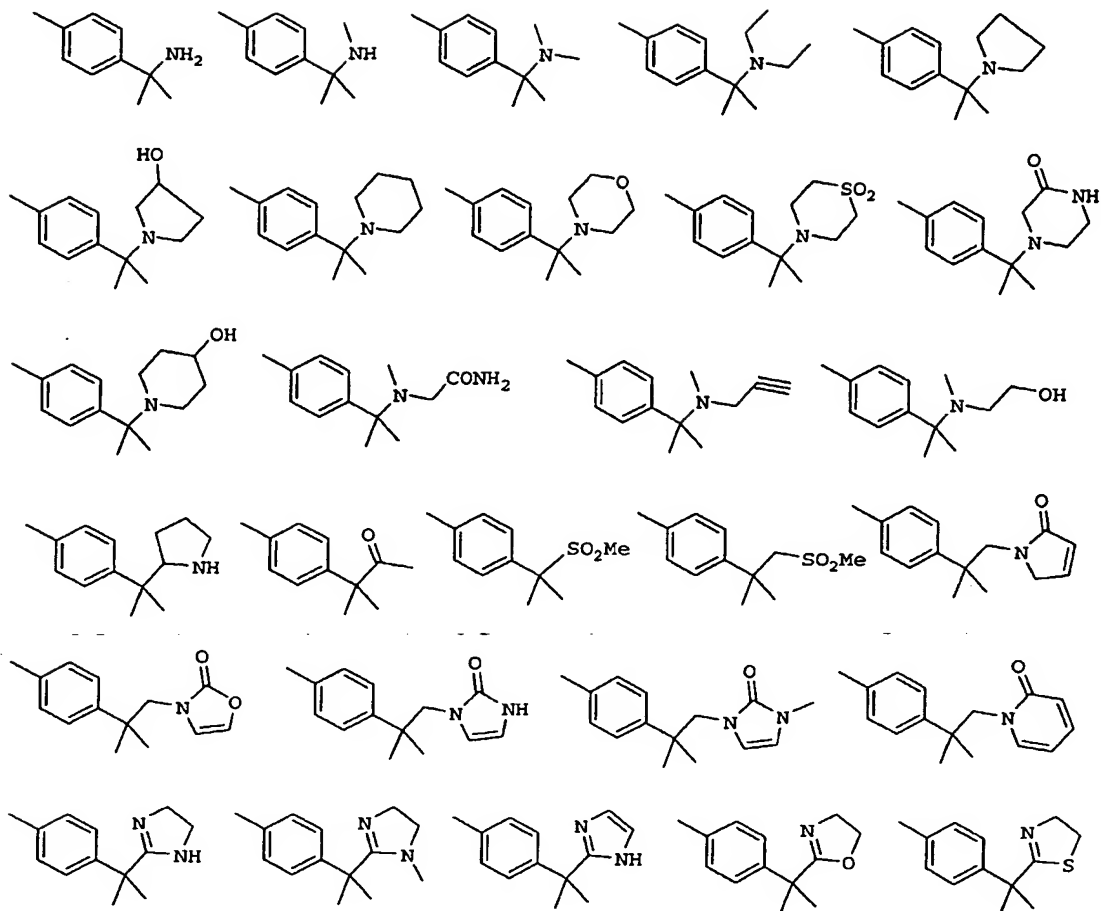


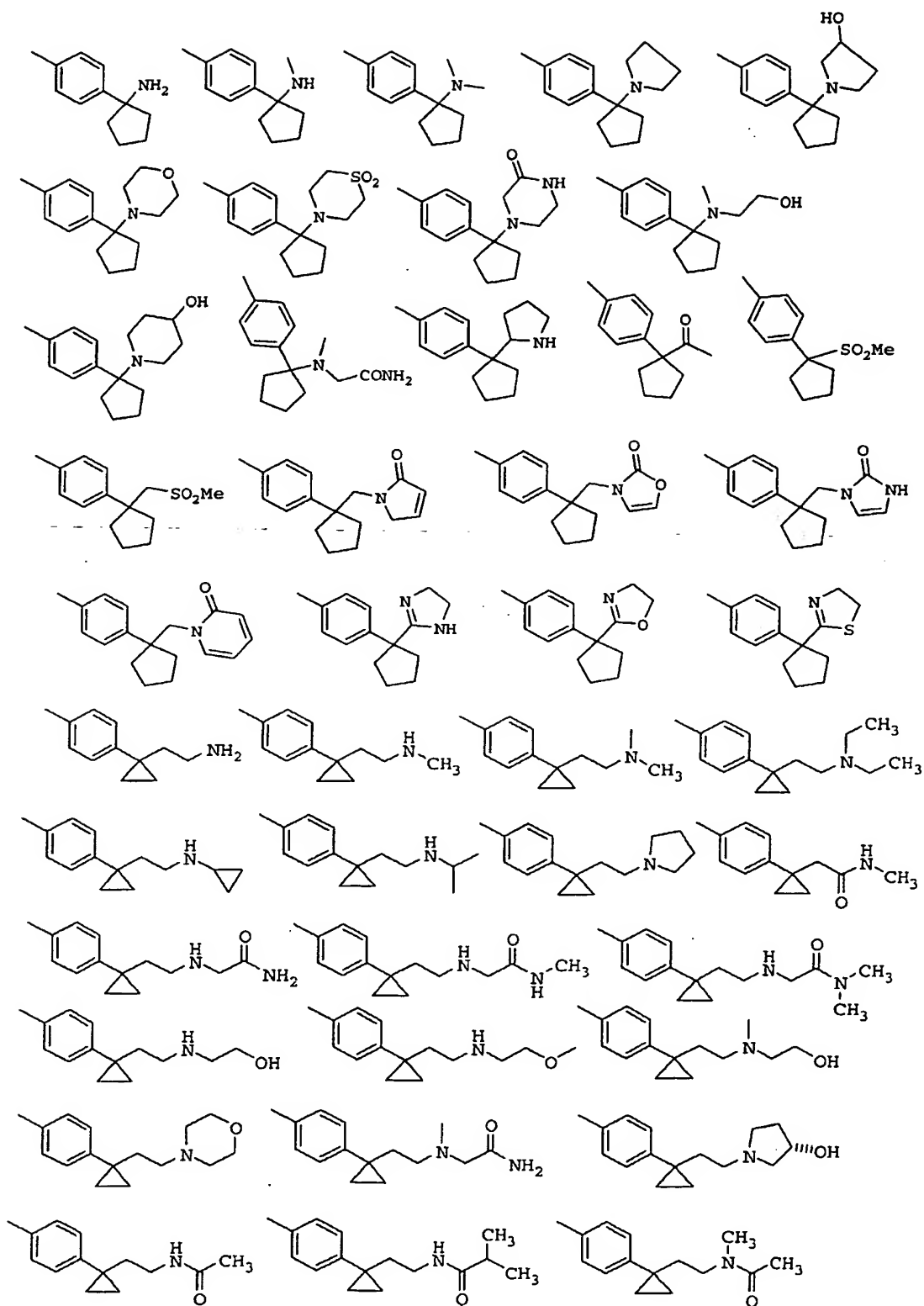


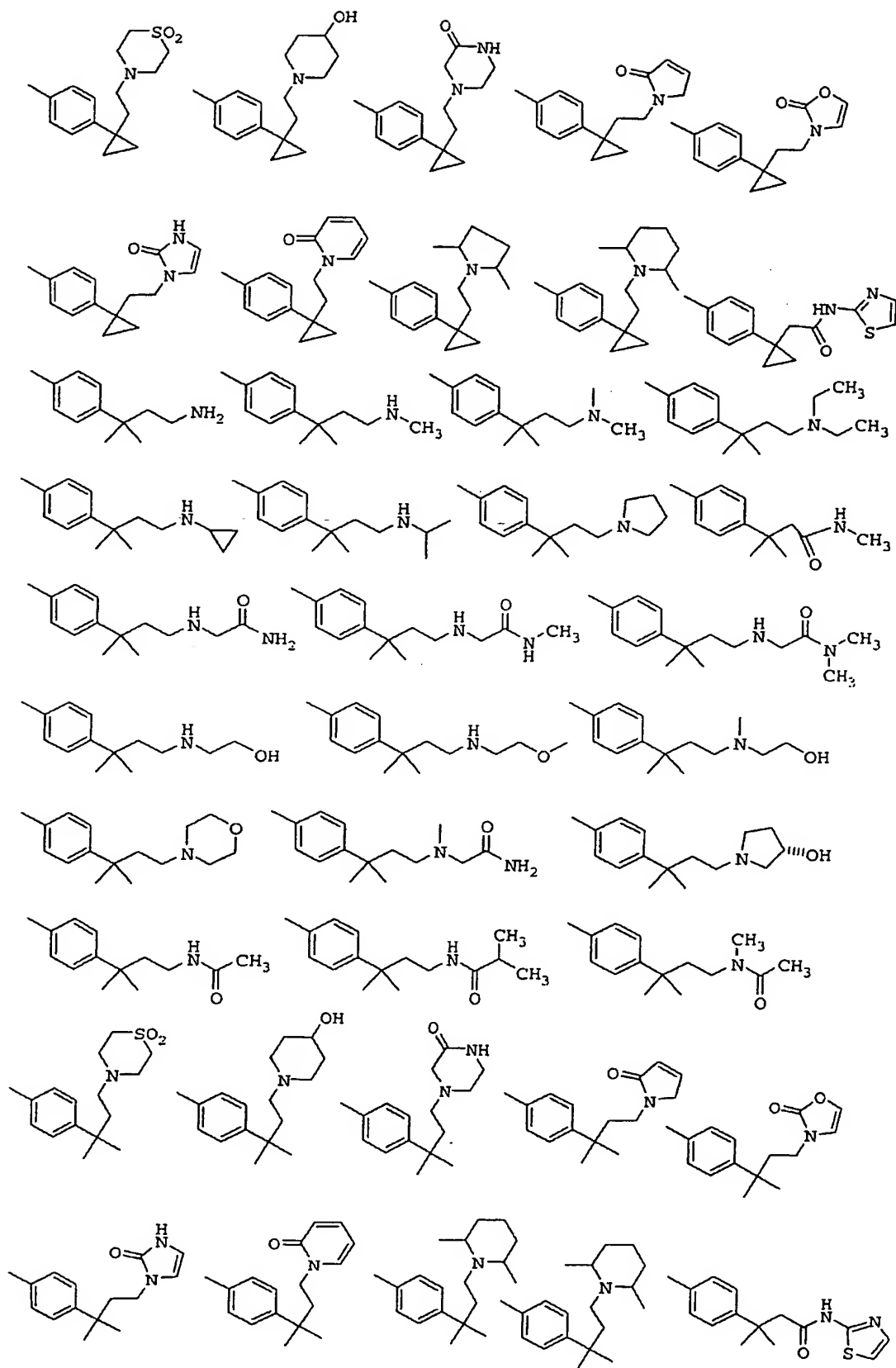


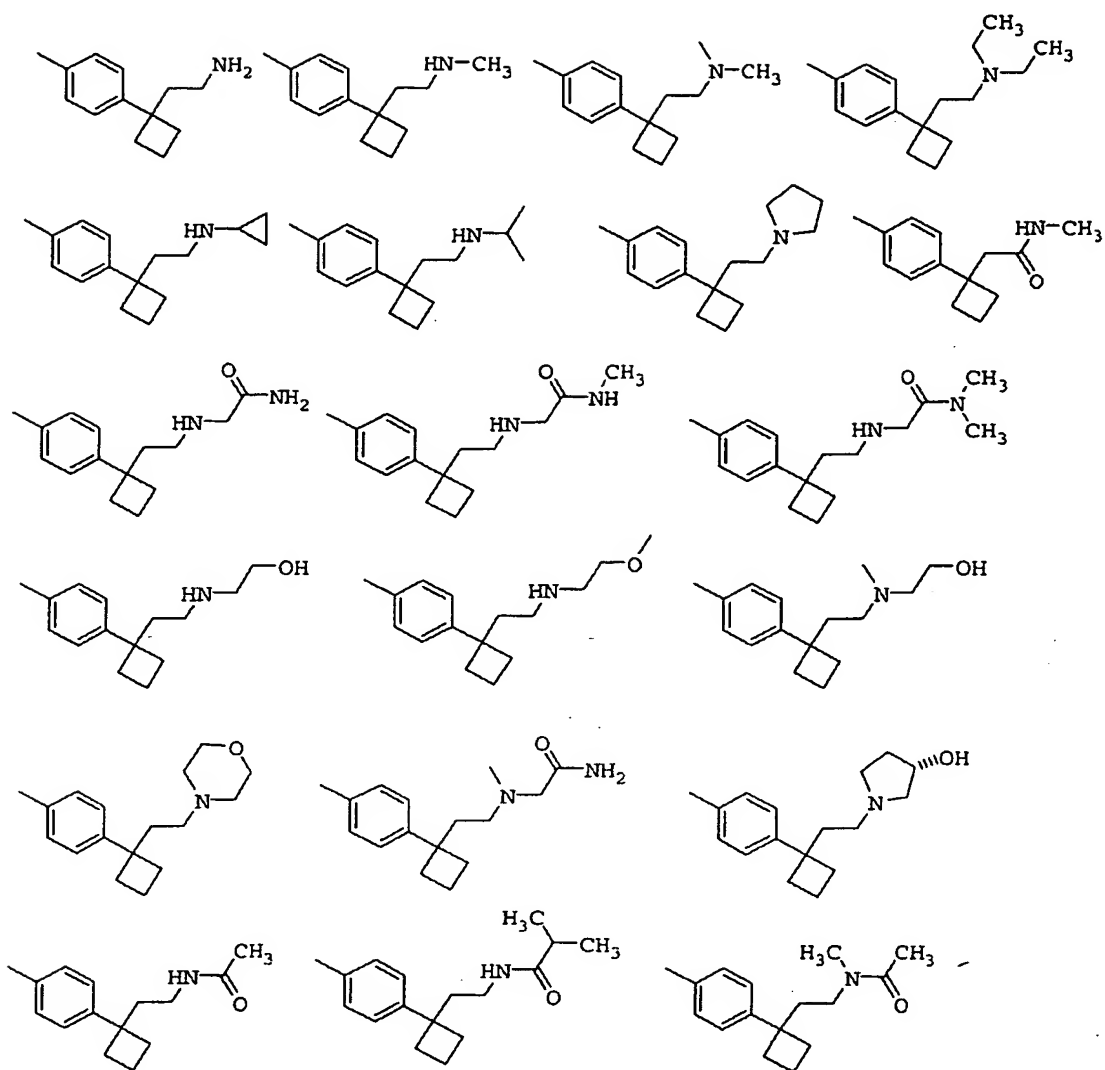


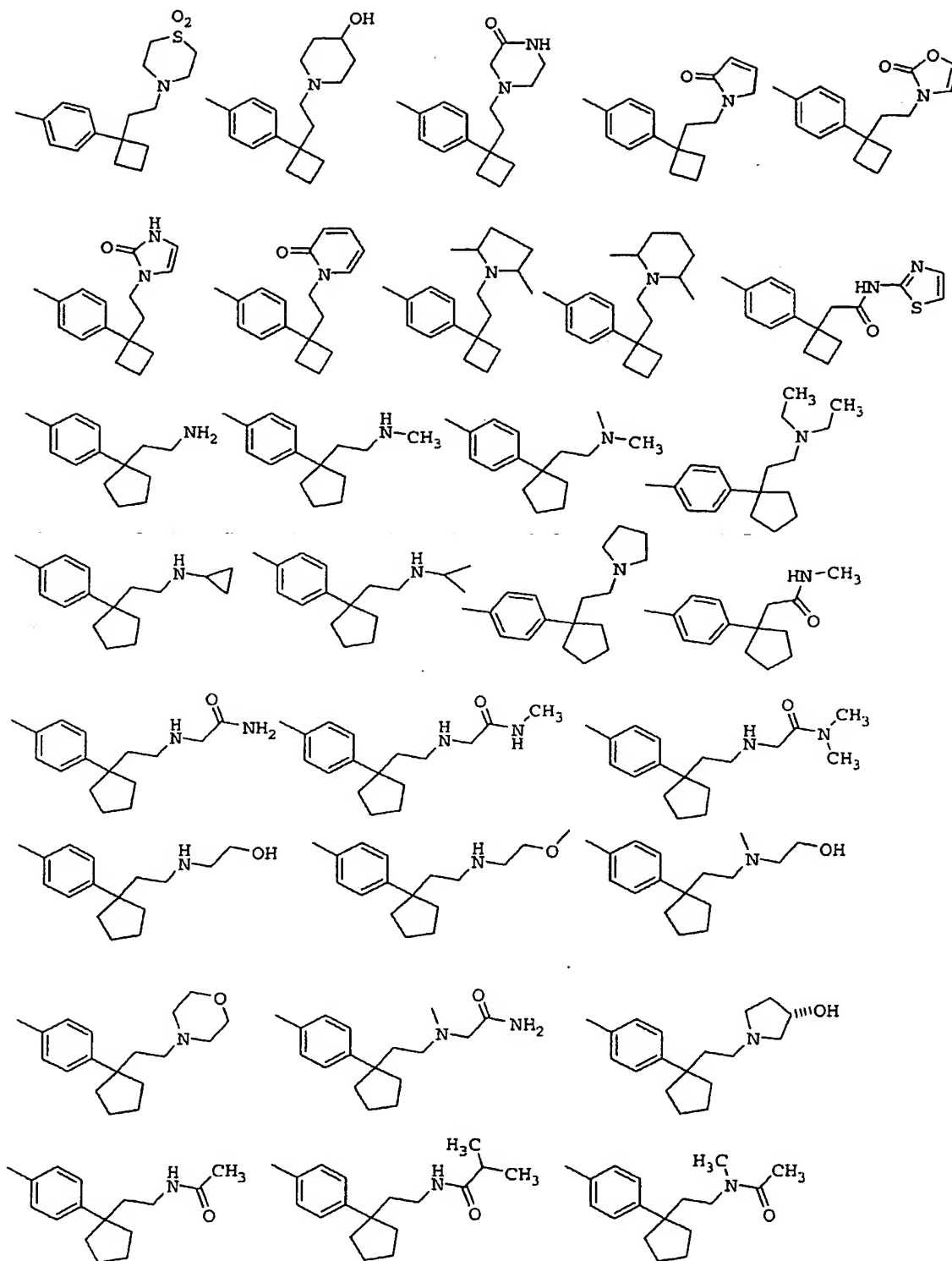


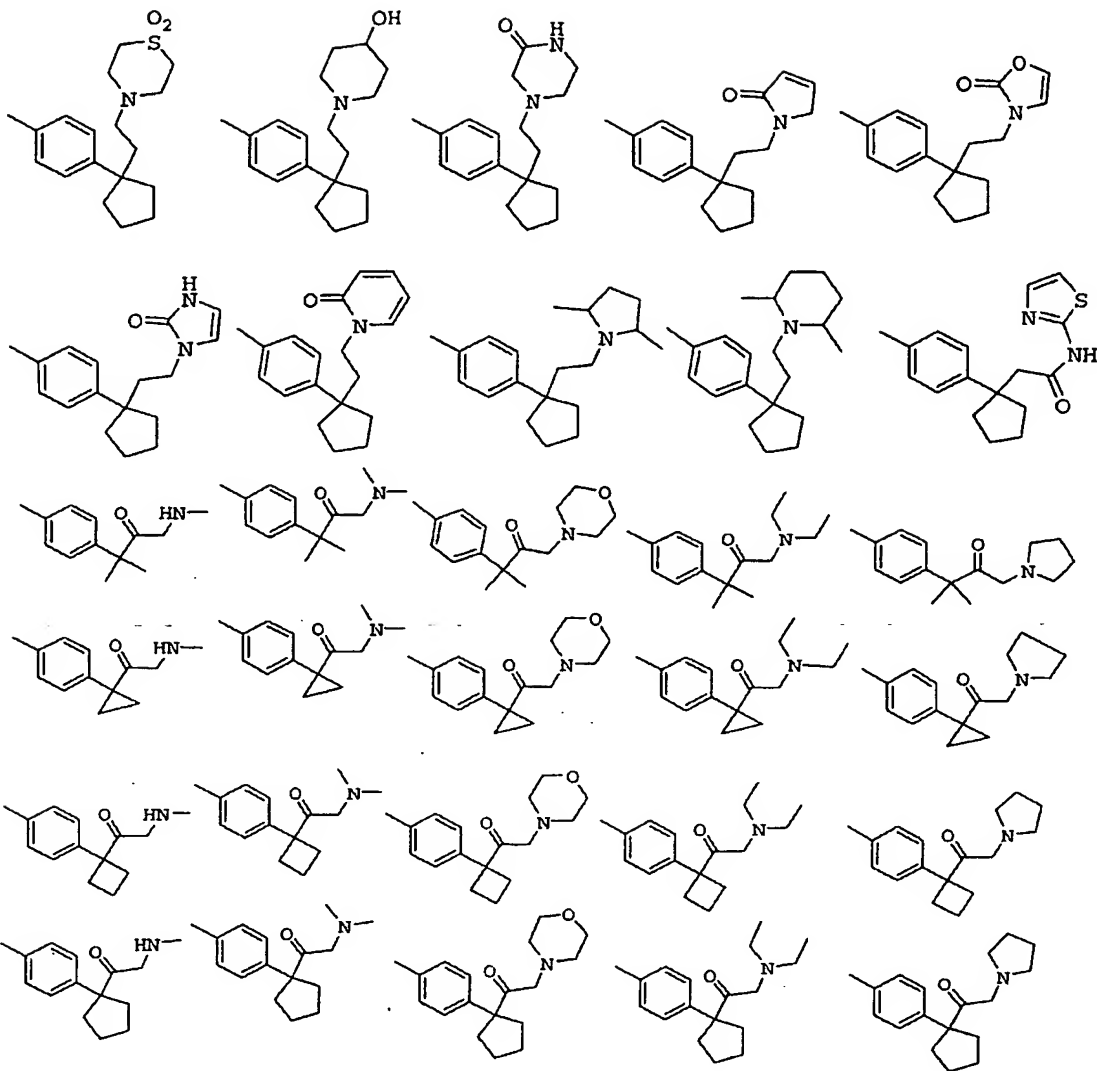












5

15. A compound according to Claim 1, wherein the compound is selected from the group:

1-[(6-chloro-2-naphthyl)sulfonyl]-4-{4-[1-(1-
 10 pyrrolidinylmethyl)cyclopropyl]benzoyl}piperazine;
 5-chloro-N-(5-chloro-2-pyridinyl)-2-({4-[1-(1-
 pyrrolidinylmethyl)cyclopropyl]benzoyl}amino)benzamide
 ;

15

- N*-{4-[1,1-dimethyl-2-(1-pyrrolidinyl)ethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 5 *N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*⁵-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-pyrazole-3,5-dicarboxamide;
- 10 3-cyano-*N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-pyrazole-5-carboxamide;
- N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-3-(trifluoromethyl)-1*H*-pyrazole-5-
- 15 carboxamide;
- N*-{4-[2-(dimethylamino)-1,1-dimethylethyl]phenyl}-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 20 *N*-(4-{1-[(dimethylamino)methyl]cyclopentyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*-(4-{1-[(dimethylamino)methyl]cyclobutyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 25 *N*-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 1-(2,3-dihydro-1*H*-indol-6-yl)-*N*⁵-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1*H*-pyrazole-
- 30 3,5-dicarboxamide;
- 1-(2,3-dihydro-1*H*-indol-6-yl)-*N*⁵-(4-{1-[(dimethylamino)methyl]cyclopropyl}phenyl)-1*H*-
- 35 pyrazole-3,5-dicarboxamide;

5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[2-(dimethylamino)-
1,1-dimethylethyl]benzoyl}amino)benzamide;

5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(methylamino)methyl]cyclopropyl}benzoyl)amino]benzami
de;

10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
(methoxymethyl)cyclopropyl]benzoyl}amino)benzamide;

15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(dimethylamino)methyl]cyclopropyl}benzoyl)amino]benza
mide;

5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-methyl-1-
pyrrolidinyl)methyl]cyclopropyl}benzoyl)amino]benzamid
e;

20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-oxo-1-
pyrrolidinyl)methyl]cyclopropyl}benzoyl)amino]benzamid
e;

25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(isopropylamino)methyl]cyclopropyl}benzoyl)amino]benz
amide;

30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(cyclopropylamino)methyl]cyclopropyl}benzoyl)amino]be
nzamide;

35 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[(cyclobutylamino)methyl]cyclopropyl}benzoyl)amino]ben
zamide;

- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-hydroxyethyl)amino)methyl}cyclopropyl)benzoyl}amino}benzamide;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-hydroxyethyl)(methyl)amino)methyl}cyclopropyl)benzoyl}amino}benzamide;
- 10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[(3-hydroxy-1-pyrrolidinyl)methyl]cyclopropyl)benzoyl}amino}benzamide;
- 15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[(4-hydroxy-1-piperidinyl)methyl]cyclopropyl)benzoyl}amino}benzamide;
- 20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[(2-oxo-1-piperidinyl)methyl]cyclopropyl)benzoyl}amino}benzamide;
- 25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[(2-oxo-1-imidazolidinyl)methyl]cyclopropyl)benzoyl}amino}benzamide;
- 30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl)benzoyl}amino}benzamide;
- 2-{{4-(1-{{[acetyl(methyl)amino)methyl}cyclopropyl)benzoyl}amino}-5-chloro-*N*-(5-chloro-2-pyridinyl)benzamide;

- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
({methyl (methylamino) carbonyl} amino) methyl} cyclopropyl
1) benzyl} amino) benzamide;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-
{ {methyl (methylsulfonyl) amino} methyl} cyclopropyl) benzy
1} amino} benzamide;
- 10 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-
[{methylsulfonyl} amino] cyclopropyl) benzyl} amino] benzam
ide;
- 15 2-({4-[1-(acetylamino) cyclopropyl] benzyl} amino)-5-chloro-*N*-
(5-chloro-2-pyridinyl) benzamide;
- 20 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-
hydroxyethyl) amino} methyl} cyclopropyl) benzyl} amino} ben
zamide;
- 25 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-{{(2-
hydroxyethyl) (methyl) amino} methyl} cyclopropyl) benzyl} a
mino} benzamide;
- 30 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(1,3-thiazol-2-
ylamino) methyl] cyclopropyl) benzoyl} amino] benzamide;
- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-[(4-{1-[(2-methyl-1*H*-
imidazol-1-
yl) methyl] cyclopropyl) benzoyl} amino] benzamide;
- 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-({4-[1-
({ {methylamino) carbonyl} amino} methyl} cyclopropyl) benz
oyl} amino) benzamide;

- methyl [1-(4-{{(4-chloro-2-{{(5-chloro-2-pyridinyl) amino} carbonyl} phenyl) amino} carbonyl} phenyl) cyclopropyl] methylcarbamate;
- 5 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-
{{(methylsulfonyl) amino} methyl} cyclopropyl) benzoyl} amino} benzamide;
- 10 2-{{4-[1-(2-amino-2-oxoethyl) cyclopropyl] benzoyl} amino)-5-chloro-*N*-(5-chloro-2-pyridinyl) benzamide;
- 15 5-chloro-*N*-(5-chloro-2-pyridinyl)-2-{{4-(1-[2-(dimethylamino)-2-oxoethyl] cyclopropyl) benzyl} amino} benzamide;
- 20 *N*-(4-[1-(2-amino-2-oxoethyl) cyclopropyl] phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- N*-(4-[1-(aminomethyl) cyclopropyl] phenyl)-1-(4-methoxyphenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 25 1-(4-methoxyphenyl)-*N*-(4-{1-
[(methylamino) methyl] cyclopropyl} phenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 30 1-(4-methoxyphenyl)-*N*-(4-[1-(1-pyrrolidinylmethyl) cyclopropyl] phenyl)-1*H*-1,2,3-triazole-5-carboxamide;
- 35 1-(4-methoxyphenyl)-*N*⁵-(4-[1-(1-pyrrolidinylmethyl) cyclopropyl] phenyl)-1*H*-pyrazole-3,5-dicarboxamide;

- 1-(4-methoxyphenyl)-N⁵-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1H-pyrazole-3,5-dicarboxamide;
- 5 1-(4-methoxyphenyl)-N⁵-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-1H-pyrazole-3,5-dicarboxamide;
- 10 3-cyano-1-(4-methoxyphenyl)-N-(4-{1-[(methylamino)methyl]cyclopropyl}phenyl)-1H-pyrazole-5-carboxamide;
- 15 3-cyano-1-(4-methoxyphenyl)-N-{4-[1-(1-pyrrolidinylmethyl)cyclopropyl]phenyl}-1H-pyrazole-5-carboxamide;
- 20 3-cyano-1-(4-methoxyphenyl)-N-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1H-pyrazole-5-carboxamide;
- 1-(4-methoxyphenyl)-3-(methylsulfonyl)-N-(4-{1-[(2-oxo-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1H-pyrazole-5-carboxamide;
- 25 N-(4-{1-[(3-hydroxy-1-pyrrolidinyl)methyl]cyclopropyl}phenyl)-1-(4-methoxyphenyl)-3-(methylsulfonyl)-1H-pyrazole-5-carboxamide;
- 30 5-chloro-thiophene-2-carboxylic acid {1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;

- 3-chloro-1H-indole-6-carboxylic acid {1-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid {1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoyl]-pyrrolidin-3-yl}-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 15 2{4-[4-chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;
- 20 2{4-[4-chloro-2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propyl alcohol;
- 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-[2-(ethylamino)-1,1-dimethylethyl]benzoyl}amino)benzamide;
- 25 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl}amino)benzamide;
- 5-chloro-N-(5-chloropyridin-2-yl)-2-({4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl}amino)benzamide;
- 30 2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;
- 35 2-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-4-methoxy-phenylcarbamoyl]-phenyl}-2-methyl-propionic acid methyl ester;

- N*-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)benzamide;
- 5 *N*-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzoyl]amino)-5-methoxybenzamide;
- N*-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)benzamide;
- 10 *N*-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-morpholin-4-ylethyl)benzoyl]amino)benzamide;
- N*-(5-chloropyridin-2-yl)-2-([4-(1,1-dimethyl-2-pyrrolidin-1-ylethyl)benzoyl]amino)-5-methoxybenzamide;
- 15 2-([4-{2-[acetyl(methyl)amino]-1,1-dimethylethyl)benzoyl]amino)-*N*-(5-chloropyridin-2-yl)benzamide;
- 20 2-(4-{[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylamino]methyl}-phenyl)-2-methyl-propionic acid methyl ester;
- 25 5-chloro-*N*-(5-chloropyridin-2-yl)-2-([4-(2-hydroxy-1,1-dimethylethyl)benzyl]amino)benzamide;
- 5-chloro-*N*-(5-chloro-pyridin-2-yl)-2-[4-(2-dimethylamino-1,1-dimethyl-ethyl)-benzylamino]-benzamide;
- 30 *N*-(5-chloropyridin-2-yl)-2-([4-[1-(hydroxymethyl)cyclopropyl]benzoyl]amino)-5-methoxybenzamide;
- 35 *N*-(5-chloropyridin-2-yl)-5-methoxy-2-([4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl]amino)benzamide;

N-(5-chloropyridin-2-yl)-2-((4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]benzoyl)amino)benzamide;

5 1-{4-[2-(5-chloro-pyridin-2-ylcarbamoyl)-phenylcarbamoyl]-phenyl}-cyclopropanecarboxylic acid methyl ester;

N-(5-chloropyridin-2-yl)-2-((4-[1-(hydroxymethyl)cyclopropyl]benzoyl)amino)benzamide;

10

6-chloro-3-(5-chloropyridin-2-yl)-2-[4-(1,1-dimethyl-2-morpholin-4-ylethyl)phenyl]quinazolin-4(3*H*)-one;

15 3-(5-chloropyridin-2-yl)-2-{4-[1-(pyrrolidin-1-ylmethyl)cyclopropyl]phenyl}quinazolin-4(3*H*)-one;

2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-methylamino-ethyl)-cyclopropyl]-phenyl}-amide;

20

2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;

25 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid {4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclopropyl]-phenyl}-amide;

30 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid [4-(1-{2-[(2-hydroxy-ethyl)-methylamino]-ethyl}-cyclopropyl)-phenyl]-amide;

35 2-(4-methoxy-phenyl)-5-trifluoromethyl-2*H*-pyrazole-3-carboxylic acid (4-{1-[2-(carbamoylmethyl-methylamino)-ethyl]-cyclopropyl}-phenyl)-amide;

- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-morpholin-4-yl-ethyl)-cyclopropyl]-phenyl}-amide;
- 5 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 10 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 15 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclobutyl)-phenyl]-amide;
- 20 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [4-(1-carbamoylmethyl-cyclobutyl)-phenyl]-amide;
- 25 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-methylamino-ethyl)-cyclobutyl]-phenyl}-amide;
- 30 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-pyrrolidin-1-yl-ethyl)-cyclobutyl]-phenyl}-amide;
- 35 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-morpholin-4-yl-ethyl)-cyclobutyl]-phenyl}-amide;

- 2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopentyl]-phenyl}-amide;
- 5 5-cyano-2-(4-methoxy-phenyl)-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 10 2-(4-methoxy-phenyl)-5-methyl-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 15 1-(4-methoxy-phenyl)-1H-pyrazole-3,5-dicarboxylic acid 3-amide 5-({4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide);
- 20 5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazole-3-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid {4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-phenyl}-amide;
- 25 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid [4-(1-carbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 30 3-(4-methoxy-phenyl)-3H-[1,2,3]triazole-4-carboxylic acid [4-(1-methylcarbamoylmethyl-cyclopropyl)-phenyl]-amide;
- 2-[1-(4-{2-[3-(4-methoxy-phenyl)-3H-[1,2,3]triazol-4-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-N-methyl-acetamide;
- 35 2-[1-(4-{2-[3-(4-methoxy-phenyl)-3H-[1,2,3]triazol-4-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-acetamide;

2-[1-(4-{2-[2-(4-methoxy-phenyl)-5-trifluoromethyl-2H-pyrazol-3-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-acetamide;

5

2-[1-(4-{2-[5-cyano-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-acetamide;

10

2-[1-(4-{2-[5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-acetamide;

15

2-[1-(4-{2-[5-methanesulfonyl-2-(4-methoxy-phenyl)-2H-pyrazol-3-yl]-2-oxo-ethyl}-phenyl)-cyclopropyl]-N-methyl-acetamide;

20

5-chloro-N-(5-chloro-2-pyridinyl)-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

N-(5-chloro-2-pyridinyl)-5-methoxy-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

25

N-(5-chloro-2-pyridinyl)-5-fluoro-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

30

N-(5-chloro-2-pyridinyl)-5-methyl-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

35

N-(5-chloro-2-pyridinyl)-5-methylsulfonyl-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

N-(5-chloro-2-pyridinyl)-5-cyano-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

5 N-(5-chloro-2-pyridinyl)-2-({4-[1-(2-dimethylamino-ethyl)cyclopropyl]benzoyl}amino)benzamide;

3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-pyridine-2-carboxylic acid (5-chloro-pyridin-2-yl)-amide;

10

N-(5-chloro-pyridin-2-yl)-4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-nicotinamide;

15 N-(5-chloro-pyridin-2-yl)-3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-isonicotinamide;

N-(5-chloro-pyridin-2-yl)-2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-nicotinamide;

20 5-chloro-N-(5-chloro-2-pyridinyl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

25 N-(5-chloro-2-pyridinyl)-5-methoxy-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

30 N-(5-chloro-2-pyridinyl)-5-fluoro-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

N-(5-chloro-2-pyridinyl)-5-methyl-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;

35

- N-(5-chloro-2-pyridinyl)-5-methylsulfonyl-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}benzoylamino)benzamide;
- 5 N-(5-chloro-2-pyridinyl)-5-cyano-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;
- 10 N-(5-chloro-2-pyridinyl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)benzamide;
- 3-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-pyridine-2-carboxylic acid (5-chloro-pyridin-2-yl)-amide;
- 15 N-(5-chloro-pyridin-2-yl)-4-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-nicotinamide;
- 20 N-(5-chloro-pyridin-2-yl)-3-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-isonicotinamide;
- N-(5-chloro-pyridin-2-yl)-2-(4-{1-[2-(2-oxo-pyrrolidin-1-yl)-ethyl]-cyclopropyl}-benzoylamino)-nicotinamide;
- 25 3-chloro-1H-indole-6-carboxylic acid {4-dimethylcarbamoyl-2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid {5-dimethylcarbamoyl-2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide;
- 35 3-chloro-1H-indole-6-carboxylic acid {4-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-tetrahydro-pyran-3-yl}-amide;

- 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-tetrahydro-pyran-4-yl}-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid {1,1-dioxo-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-hexahydro-1 λ ⁶-thiopyran-4-yl}-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid {1,1-dioxo-4-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-hexahydro-1 λ ⁶-thiopyran-3-yl}-amide;
- 15 3-chloro-1H-indole-6-carboxylic acid {1-acetyl-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl}-amide;
- 20 3-chloro-1H-indole-6-carboxylic acid {1-acetyl-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl}-amide;
- 4-[(3-chloro-1H-indole-6-carbonyl)-amino]-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidine-1-carboxylic acid methyl ester;
- 25 3-chloro-1H-indole-6-carboxylic acid {1-(2-methoxy-acetyl)-3-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-piperidin-4-yl}-amide;
- 30 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-tetrahydro-furan-3-yl}-amide;

- 5-chloro-thiophene-2-carboxylic acid {1-acetyl-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidin-3-yl}-amide;
- 5 5-chloro-thiophene-2-carboxylic acid {1-cyclopropanecarbonyl-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidin-3-yl}-amide;
- 10 3-[(5-chloro-thiophene-2-carbonyl)-amino]-4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-pyrrolidine-1-carboxylic acid methyl ester;
- 15 5-chloro-thiophene-2-carboxylic acid [4-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1-(2-methoxy-acetyl)-pyrrolidin-3-yl]-amide;
- 20 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-4-dimethylcarbamoyl-cyclopentyl}-amide;
- 25 5-chloro-thiophene-2-carboxylic acid {1-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-indan-2-yl}-amide;
- 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1,2,3,4-tetrahydro-naphthalen-2-yl}-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid {3-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-7-oxa-bicyclo[2.2.1]hept-2-yl}-amide;
- 35 5-chloro-thiophene-2-carboxylic acid {2-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-4-dimethylcarbamoyl-cyclopentyl}-amide;

- 5-chloro-thiophene-2-carboxylic acid {8-[4-(1-dimethylaminomethyl-cyclopropyl)-benzoylamino]-1-oxa-spiro[4.4]non-7-yl}-amide;
- 5 5-chloro-thiophene-2-carboxylic acid (8-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1-oxa-spiro[4.4]non-7-yl)-amide;
- 10 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-cyclopentyl)-amide;
- 15 5-chloro-thiophene-2-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-4-dimethylcarbamoyl-cyclopentyl)-amide;
- 20 3-[(5-chloro-thiophene-2-carbonyl)-amino]-4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-pyrrolidine-1-carboxylic acid methyl ester;
- 25 5-chloro-thiophene-2-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-furan-3-yl)-amide;
- 30 3-chloro-1H-indole-6-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-cyclohexyl)-amide;
- 35 3-chloro-1H-indole-6-carboxylic acid (2-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-4-dimethylcarbamoyl-cyclohexyl)-amide;
- 4-[(3-Chloro-1H-indole-6-carbonyl)-amino]-3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-piperidine-1-carboxylic acid methyl ester;

- 3-chloro-1H-indole-6-carboxylic acid (3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1,1-dioxo-hexahydro-1 λ ⁶-thiopyran-4-yl)-amide;
- 5 3-chloro-1H-indole-6-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-1,1-dioxo-hexahydro-1 λ ⁶-thiopyran-3-yl)-amide;
- 10 3-chloro-1H-indole-6-carboxylic acid (4-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-pyran-3-yl)-amide;
- 15 3-chloro-1H-indole-6-carboxylic acid (3-{4-[1-(2-dimethylamino-ethyl)-cyclopropyl]-benzoylamino}-tetrahydro-pyran-4-yl)-amide;
- (1R, 2S)-5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide;
- 20 (1R, 2S)-3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclopentyl}-amide;
- 25 (1R, 2S)-5-chloro-thiophene-2-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-benzoylamino]-cyclohexyl}-amide; and,
- 30 Cis-3-chloro-1H-indole-6-carboxylic acid {2-[4-(1-pyrrolidin-1-ylmethyl-cyclopropyl)-phenylcarbamoyl]-cyclohexyl}-amide;

or a pharmaceutically acceptable salt form thereof.

16. A pharmaceutical composition, comprising: a
pharmaceutically acceptable carrier and a therapeutically
effective amount of a compound of Claim 1, 2, 3, 4, 5, 6,
7, 8, 9, 10, 11, 12, 13, 14, or 15 or a pharmaceutically
5 acceptable salt thereof.

17. A compound of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10,
11, 12, 13, 14, or 15 for use in therapy.
10

18. Use of a compound of Claim 1, 2, 3, 4, 5, 6, 7, 8,
9, 10, 11, 12, 13, 14, or 15 for the manufacture of a
medicament for the treatment of a thromboembolic disorder.
15

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US03/13893

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 31/41, 3144, 31/435; C07D 213/14, 213/75, 471/04

US CL : 514/300, 303, 352, 406, 407; 546/117, 119, 309; 548/364.7, 369.4, 369.7

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/300, 303, 352, 406, 407; 546/117, 119, 309; 548/364.7, 369.4, 369.7

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CAS ONLINE- Structure searches**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A,P	US 6,456,656 B2 (ZHOU et al) 15 October 2002.	1-18

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

14 July 2003 (14.07.2003)

Date of mailing of the international search report

08 AUG 2003

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US
Commissioner for Patents
P.O. Box 1450
Alexandria, Virginia 22313-1450

Facsimile No. (703)305-3230

Authorized officer

Bernard Dentz

Telephone No. 703 308-1235

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

I. Claims 1-8 and 16-18 drawn to pyrazolo and triazolopyridines.

II. Claims 1 and 9-18 drawn to pyrazoles.

III. Claims 1 and 9-18 drawn to compounds where the M ring is benzene. See claim 14, fourth and fifth structures.

IV. Claims 1 and 9-18 drawn to cpds. where M is cyclohexane or cyclopentane. See claim 14, sixth and seventh structures

V. Claims 1 and 9-18 drawn to cpds. where M is piperidine. See claim 14, eighth and ninth structures.

In covering a multitude of different ring structures there is not a single common core. See PCT rule 13.1-13.4.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US03/13893

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claim Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐
☐

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.